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TO:

Shirley Gembeh

Art Unit:

1614

Location: REM-3A44/3C70

Serial Number: 10/712423

Monday, March 13, 2006

From: **Beverly Shears**

Location: Biotech-Chem Library

REM 1A54

Phone:

571-272-2528

beverly.shears@uspto.gov

Search Notes

Shirley,

If you have any questions regarding the search strategy, pls. contact me.

Beverly



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STR

L1

C=0 @19 20

2 7 || 11 14

1 C 3 CH2 ~ CH2 ~ G1 ~ C ~ CH2

1 C 3 CH2 ~ CH2 ~ G1 ~ C ~ CH2

1 C 6 C 4

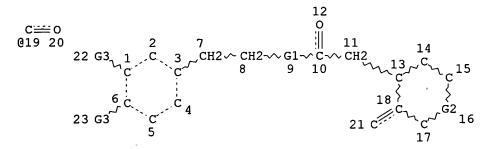
21 C C 6 C 17

VAR G1=0/S/CH2/19 VAR G2=0/S/CH2/19 NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 21

STEREO ATTRIBUTES: NONE

L2 (132) SEA FILE=REGISTRY SSS FUL L1 L3 STR



VAR G1=O/S/CH2/19
VAR G2=O/S/CH2/19
VAR G3=OH/N/S
NODE ATTRIBUTES:
CONNECT IS X2 RC AT 15
CONNECT IS M3 RC AT 17
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

L4 32 SEA FILE=REGISTRY SUB=L2 SSS FUL L3

100.0% PROCESSED 127 ITERATIONS

SEARCH TIME: 00.00.01

32 ANSWERS

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L5 540 SEA ABB=ON PLU=ON L4

L6 30 SEA ABB=ON PLU=ON L5 AND (?CANCER? OR ?CARCIN? OR

?TUMOUR? OR ?TUMOR? OR ?NEOPLAS?)

L7 13 SEA ABB=ON PLU=ON L6 NOT (PY=>2002 OR PD=>20021209)

E3 THROUGH E4 ASSIGNED

L7 ANSWER 1 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:296202 CAPLUS

DOCUMENT NUMBER: 136:293669

TITLE: Oleuropein and related degradation compounds of

olives with antioxidant activity

AUTHOR(S): Schmandke, Horst

CORPORATE SOURCE: Bergholz-Rehbrucke, 14558, Germany

SOURCE: Ernaehrungs-Umschau (2001), 48(12), 490-492

CODEN: ERUMAT; ISSN: 0014-021X

PUBLISHER: Umschau Zeitschriftenverlag Breidenstein

DOCUMENT TYPE: Journal; General Review

LANGUAGE: German

AB A review. Fresh olives contain the bitter principle oleuropein, its aglycon and the degradation products elenolic acid glucoside, hydroxytyrosol and its glucosides. During the debittering of olives by Spanish-, Californian- or Greek-style processing oleuropein further on is degradated to the earlier mentioned compds. who - together with the residual oleuropein - finally are being removed from the olives into the processing brine by diffusion. Non raffinated olive oils (extra virgin and virgin oils) contain beside small contents of oleuropein first of all his aglycon, the corresponding dialdehydic form (oleuropeindial), small concns. of decarbomethoxyoleuropeindial and hydroxytyrosol. By oil raffination all phenolic compds. are eliminated vastly. Oleuropein is poorly and hydroxytyrosol dose-dependently absorbed. They are excreted urinarily as glucuronide conjugates of hydroxytyrosol and 4-hydroxy-3-methoxy-phenethanol. In vitro studies demonstrated that oleuropein and hydroxytyrosol have antioxidant activities in cell-free model solns., in LDL, platelets, leukocytes, neutrophils and epithelial intestinal cells. This antioxidant effect inhibits the arachidonic acid metabolism by inhibition of the 5- and 12-lipoxygenase activities and the formation of 5- and 12-hydroxy-5,8,10,14-eicosatetraenoic acid, F2-isoprostane, leukotriene B4 and thromboxane B2. The platelet aggregation is reduced. These facts are confirmed by a dose-dependent decrease of urinary excretion of F2-isoprostane in humans. In vitro oleuropein shows cytostatic activity against McCoy cells, indicating an antitumoral capacity. Oleuropein inhibits cytochrome P 450 and can therefore effect the bioavailability of drugs.

IT 32619-42-4, Oleuropein

RL: BSU (Biological study, unclassified); BIOL (Biological study) (oleuropein and related degradation compds. of olives with antioxidant activity)

RN 32619-42-4 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S,3E,4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

REFERENCE COUNT:

THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 2 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

8

2001:818973 CAPLUS ACCESSION NUMBER:

137:15715 DOCUMENT NUMBER:

TITLE: The inhibitory effects of compounds from olive

leaf on tumor necrosis factor production

and on β -hexosaminidase release

AUTHOR(S): Nishibe, Sansei; Han, Yingmei; Noguchi, Yukari;

Ueda, Hiroshi; Yamazaki, Masatoshi; Mizutani, Kenji; Kambara, Toshimitsu; Kishida, Naoko

Faculty of Pharmaceutical Sciences, Health CORPORATE SOURCE:

Sciences University of Hokkaido, Ishikari-Tobetsu,

Hokkaido, 061-0293, Japan

Natural Medicines (Tokyo, Japan) (2001), 55(4), SOURCE:

205-208

CODEN: NMEDEO; ISSN: 1340-3443 Japanese Society of Pharmacognosy

PUBLISHER: DOCUMENT TYPE: Journal

English LANGUAGE:

The extraction and isolation of olive leaf gave luteolin 7-0-glucoside, luteolin 4'-O-glucoside and oleuropein as the major components. inhibitory effects of these compds. on tumor necrosis factor $(TNF-\alpha)$ production and on β -hexosaminidase release from rat basophilic leukemia (RBL-2H3) cells, which were both recently found to be linked to allergic reaction, were examined Oleuropein showed a potent inhibitory effect on $TNF-\alpha$ production Luteolin 4'-O-glucoside showed a strong inhibitory effect on β -hexosaminidase release (IC50:17.1 μ g/mL).

32619-42-4P, Oleuropein

RL: NPO (Natural product occurrence); PAC (Pharmacological activity); PUR (Purification or recovery); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); PREP (Preparation); USES (Uses)

> Shears 571-272-2528 Searcher :

(inhibitory effects of compds. from olive leaf on tumor necrosis factor production and on β -hexosaminidase release)

32619-42-4 CAPLUS RN

2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-CN 3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S, 3E, 4S) - (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

REFERENCE COUNT:

THERE ARE 17 CITED REFERENCES AVAILABLE FOR 17 THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

CAPLUS COPYRIGHT 2006 ACS on STN ANSWER 3 OF 13

ACCESSION NUMBER:

2001:490007 CAPLUS

DOCUMENT NUMBER:

135:226053

TITLE:

Characterization of Phenolic Compounds in Virgin Olive Oil and Their Effect on the Formation of Carcinogenic/Mutagenic Heterocyclic Amines

in a Model System

AUTHOR(S):

Monti, Simona M.; Ritieni, Alberto; Sacchi, Raffaele; Skog, Kerstin; Borgen, Eva; Fogliano,

Vincenzo

CORPORATE SOURCE:

Dipartimento di Scienza degli Alimenti, Universita

di Napoli Federico II, Naples, 80055, Italy

SOURCE:

Journal of Agricultural and Food Chemistry (2001),

49(8), 3969-3975

:

CODEN: JAFCAU; ISSN: 0021-8561

PUBLISHER:

American Chemical Society

DOCUMENT TYPE:

Journal English

LANGUAGE:

Mutagenic heterocyclic amines (HAs) are formed at low levels during

Searcher

Shears

571-272-2528

cooking of meat and fish, and some of them are considered to be possible human carcinogens. The formation of HAs may be affected by the presence of synthetic or naturally occurring antioxidants. In the present study the effect of virgin olive oil (VOO) phenolic compds., identified and quantified by LC-MS, on the formation of HAs in a model system was evaluated. An aqueous solution of creatinine, glucose, and glycine was heated in the presence of 2 samples of VOO differing only in the composition of phenolic compds. addition of VOO to the model system inhibited the formation of 2-amino-3-methylimidazo[4,5-f]quinoxaline (IQx), 2-amino-3,8dimethylimidazo[4,5-f]quinoxaline (MeIQx), and 2-amino-3,7,8trimethylimidazo[4,5-f]quinoxaline (DiMeIQx) by between 30 and 50% compared with the control. Fresh-made olive oil, which contained a high amount of dihydroxyphenylethanol derivs., inhibited HA formation more than a 1-yr-old oil did. The inhibition of HA formation was also verified using phenolic compds. extracted from VOO.

IT 31773-95-2, Oleuropein aglycon

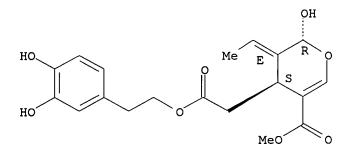
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study) (characterization of, in virgin olive oil and effect on formation of carcinogenic/mutagenic heterocyclic amines in a model system)

RN 31773-95-2 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-3,4-dihydro-2-hydroxy-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2R,3E,4S)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.



REFERENCE COUNT: 58 THERE ARE 58 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L7 ANSWER 4 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:155024 CAPLUS

DOCUMENT NUMBER: 134:310130

TITLE: Olive-oil consumption and health: the possible

role of antioxidants

AUTHOR(S): Owen, Robert W.; Giacosa, Attilio; Hull, William

E.; Haubner, Roswitha; Wurtele, Gerd; Spiegelhalder, Bertold; Bartsch, Helmut

CORPORATE SOURCE: Division of Toxicology and Cancer Risk Factors,

German Cancer Research Center, Heidelberg,

D-69120, Germany

SOURCE: Lancet Oncology (2000), 1(Oct.), 107-112

CODEN: LOANBN; ISSN: 1470-2045

PUBLISHER: Lancet Publishing Group DOCUMENT TYPE: Journal; General Review LANGUAGE: English

A review with 35 refs. In the Mediterranean basin, olive oil, along with fruits, vegetables, and fish, is an important constituent of the diet, and is considered a major factor in preserving a healthy and relatively disease-free population. Epidemiol. data show that the Mediterranean diet has significant protective effects against cancer and coronary heart disease. We present evidence that it is the unique profile of the phenolic fraction, along with high intakes of squalene and the monounsatd. fatty acid, oleic acid, which confer its health-promoting properties. The major phenolic compds. identified and quantified in olive oil belong to three different classes: simple phenols (hydroxytyrosol, tyrosol); secoiridoids (oleuropein, the aglycon of ligstroside, and their resp. decarboxylated dialdehyde derivs.); and the lignans [(+)-1-acetoxypinoresinol and (+)-pinoresinol]. All three classes have potent antioxidant properties. High consumption of extra-virgin olive oils, which are particularly rich in these phenolic antioxidants (as well as squalene and oleic acid), should afford considerable protection against cancer (colon, breast, skin), coronary heart disease, and ageing by inhibiting oxidative stress.

IT 32619-42-4, Oleuropein
RL: BAC (Biological activity or effector, except adverse); BSU

(Biological study, unclassified); BIOL (Biological study)
(Olive-oil consumption and health in relation to the possible role
of antioxidants)

RN 32619-42-4 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S,3E,4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

REFERENCE COUNT:

35 THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 5 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:906581 CAPLUS

DOCUMENT NUMBER: 134:146898

TITLE: Oleuropein, an antioxidant polyphenol from olive

oil, is poorly absorbed from isolated perfused rat

intestine

AUTHOR(S): Edgecombe, Steven C.; Stretch, Graham L.; Hayball,

Peter J.

CORPORATE SOURCE: Centre for Pharmaceutical Research, University of

South Australia, Adelaide, 5000, Australia

SOURCE: Journal of Nutrition (2000), 130(12), 2996-3002

CODEN: JONUAI; ISSN: 0022-3166

PUBLISHER: American Society for Nutritional Sciences

DOCUMENT TYPE: Journal LANGUAGE: English

AB Epidemiol. studies have shown that the incidence of heart disease and certain cancers is lower in the Mediterranean region. This has been attributed to the high consumption of olive oil in the Mediterranean diet, which contains polyphenolic compds. With antioxidant activity. Although many in vitro studies have been performed to elucidate mechanisms by which these compds. may act, there are virtually no data relating to their fate after ingestion. Therefore, we decided to investigate the intestinal absorption of one of the major olive oil polyphenolics, oleuropein. To do this, a novel in situ intestinal perfusion technique was developed, and the absorption of oleuropein was studied under both iso-osmotic and hypotonic luminal conditions. Oleuropein was absorbed, with an apparent permeability coefficient (Papp) of 1.47 ± 0.13 + 10-6 cm/s (±SE) observed under iso-osmotic conditions. The mechanism of absorption is unclear but may involve transcellular transport (SGLT1)

or paracellular movement. Under hypotonic conditions, the permeability of oleuropein was significantly greater (5.92 \pm 0.49 + 10-6 cm/s, P < 0.001). This increase is thought to be due to an increase in paracellular movement facilitated by the opening of paracellular junctions in response to hypotonicity. Overall, we determined that the olive oil polyphenolic oleuropein can be absorbed, albeit poorly, from isolated perfused rat intestine. Therefore, it is possible that it or its metabolites may confer a pos. health benefit after the consumption of olive oil, most likely via an antioxidant mechanism.

IT 32619-42-4, Oleuropein

RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)

(oleuropein, an antioxidant polyphenol from olive oil, is poorly absorbed from isolated perfused rat intestine)

RN 32619-42-4 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S,3E,4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

REFERENCE COUNT:

60 THERE ARE 60 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 6 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

2000:507306 CAPLUS

DOCUMENT NUMBER:

133:207042

TITLE:

Phenolic compounds and squalene in olive oils: the concentration and antioxidant potential of total phenols, simple phenols, secoiridoids, lignans and

squalene

AUTHOR(S): Owen, R. W.; Mier, W.; Giacosa, A.; Hull, W. E.;

Spiegelhalder, B.; Bartsch, H.

CORPORATE SOURCE: Division of Toxicology and Cancer Risk Factors,

German Cancer Research Center, Heidelberg, Germany

Food and Chemical Toxicology (2000), 38(8),

647-659

CODEN: FCTOD7; ISSN: 0278-6915

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English

SOURCE:

The aim of this study was to evaluate the phenolic antioxidant and squalene content in a range of olive and seed oils. A mean of 290 ± 38 (SEM) mg squalene/100 g was detected. However, while there was a weak significant difference between extra virgin (424 \pm 21 mg/kg) and refined virgin (340 \pm 31 mg/100 g) olive oils, highly significant differences were evident between extra virgin olive oils refined virgin olive oils and seed oils (24 \pm 5 mg/100 g). While seed oils were devoid, on average, the olive oils contained 196 ± 19 mg/kg total phenolics as judged by HPLC anal., but the value for extra virgin (232 ± 15 mg/kg) was significantly higher than that of refined virgin olive oil (62 ± 12 mg/kg). Appreciable quantities of simple phenols (hydroxytyrosol and tyrosol) were detected in olive oils, with significant differences between extra virgin (41.87 ± 6.17) and refined virgin olive oils (4.72 ± 215) . The major linked phenols were secoiridoids and lignans. Although extra virgin contained higher concns. of secoiridoids (27.72 ± 6.84) than refined olive oils (9.30 ± 3.81) this difference was not significant. On the other hand, the concentration of lignans was significantly higher in extra virgin (41.53 ± 3.93) compared to refined virgin olive oils (7.29 ± 2.56). All classes of phenolics were shown to be potent antioxidants. In future epidemiol. studies, both the nature and source of olive oil consumed should be differentiated in ascertaining cancer risk.

IT 31773-95-2, Oleuropein aglycone 32619-42-4,
 Oleuropein

RL: ANT (Analyte); BOC (Biological occurrence); BSU (Biological study, unclassified); ANST (Analytical study); BIOL (Biological study); OCCU (Occurrence)

(phenolic compds. and squalene in olive oils, concentration and antioxidant potential of total phenols, simple phenols, secoiridoids, lignans and squalene)

RN 31773-95-2 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-3,4-dihydro-2-hydroxy-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2R,3E,4S)-(9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.

RN 32619-42-4 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S,3E,4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

REFERENCE COUNT:

37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 7 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER:

2000:487644 CAPLUS

:

DOCUMENT NUMBER:

133:192309

TITLE:

Identification of lignans as major components in

the phenolic fraction of olive oil

AUTHOR(S):

Owen, Robert W.; Mier, Walter; Giacosa, Attilio; Hull, William E.; Spiegelhälder, Bertold; Bartsch,

Helmut

Searcher

Shears

571-272-2528

CORPORATE SOURCE:

Division of Toxicology and Cancer Risk Factors,

German Cancer Research Center, Heidelberg,

D-69120, Germany

SOURCE:

Clinical Chemistry (Washington, D. C.) (2000),

46(7), 976-988

CODEN: CLCHAU; ISSN: 0009-9147

PUBLISHER:

American Association for Clinical Chemistry

DOCUMENT TYPE:

Journal

English

LANGUAGE:

AB The major phenolic antioxidants in extra virgin olive oil were isolated and purified. Structural anal. was conducted using several spectroscopic techniques, including mass spectrometry and NMR. In particular, detailed 1H and 13C NMR data are presented, and several assignment errors in the literature are corrected The lignans (+)-l-acetoxypinoresinol and (+)-pinoresinol are major components of the phenolic fraction of olive oils. These lignans, which are potent antioxidants, are absent in seed oils and absent in refined virgin oils, but are present at concns. of up to 100 mg/kg (mean ± SE, $41.53 \pm 3.93 \text{ mg/kg; range, } 0.65-99.97 \text{ mg/kg)}$ in extra virgin oils. As with the simple phenols and secoiridoids, there is considerable interoil variation in lignan concns. Foods containing high amts. of lignan precursors have been found to be protective against breast, colon, and prostate cancer. Lignans, as natural components of the diet, may be important modulators of cancer chemopreventive activity.

IT 31773-95-2 32619-42-4, Oleuropein

RL: FFD (Food or feed use); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(lignan components in the phenolic fraction of olive oil)

RN 31773-95-2 CAPLUS

2H-Pyran-4-acetic acid, 3-ethylidene-3,4-dihydro-2-hydroxy-5-CN (methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2R,3E,4S)-(9CI) (CA INDEX NAME)

Absolute stereochemistry. Double bond geometry as shown.

RN 32619-42-4 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S, 3E, 4S) - (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

REFERENCE COUNT: 40 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 8 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:454807 CAPLUS

DOCUMENT NUMBER: 133:344249

TITLE: The antioxidant/anticancer potential of

phenolic compounds isolated from olive oil

AUTHOR(S): Owen, R. W.; Giacosa, A.; Hull, W. E.; Haubner,

R.; Spiegelhalder, B.; Bartsch, H.

CORPORATE SOURCE: Division of Toxicology and Cancer Risk Factors,

German Cancer Research Centre, Heidelberg,

D-69120, Germany

SOURCE: European Journal of Cancer (2000), 36(10),

1235-1247

CODEN: EJCAEL; ISSN: 0959-8049

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English

In our ongoing studies on the chemoprevention of cancer we AB have a particular interest in the health benefits of the Mediterranean diet, of which olive oil is a major component. Recent studies have shown that extravirgin olive oil contains an abundance of phenolic antioxidants including simple phenols (hydroxytyrosol, tyrosol), aldehydic secoiridoids, flavonoids and lignans (acetoxypinoresinol, pinoresinol). All of these phenolic substances are potent inhibitors of reactive oxygen species attack on, e.g., salicylic acid, 2-deoxyguanosine. Currently there is growing evidence that reactive oxygen species are involved in the etiol. of fat-related neoplasms such as cancer of the breast and colorectum. A plausible mechanism is a high intake of $\omega \text{-}6$ polyunsatd. fatty acids which are especially prone to lipid peroxidn. initiated and propagated by reactive oxygen species, leading to the formation (via α, β -unsatd. aldehydes such as

trans-4-hydroxy-2-nonenal) of highly pro-mutagenic exocyclic DNA adducts. Previous studies have shown that the colonic mucosa of cancer patients and those suffering from predisposing inflammatory conditions such as ulcerative colitis and Crohn's disease generates appreciably higher quantities of reactive oxygen species compared with normal tissue. We have extended these studies by developing accurate high performance liquid chromatog. (HPLC) methods for the quantitation of reactive oxygen species generated by the fecal The data shows that the fecal matrix supports the generation of reactive oxygen species in abundance. As yet, there is a dearth of evidence linking this capacity to actual components of the diet which may influence the colorectal milieu. However, using the newly developed methodol. we can demonstrate that the antioxidant phenolic compds. present in olive oil are potent inhibitors of free radical generation by the fecal matrix. This indicates that the study of the inter-relation between reactive oxygen species and dietary antioxidants is an area of great promise for elucidating mechanisms of colorectal carcinogenesis and possible future chemopreventive strategies.

IT 32619-42-4P

CN

RL: ANT (Analyte); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); PUR (Purification or recovery); THU (Therapeutic use); ANST (Analytical study); BIOL (Biological study); PREP (Preparation); USES (Uses)

(antioxidant/anticancer potential of phenolic compds.

isolated from olive oil)

RN 32619-42-4 CAPLUS

2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S,3E,4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

IT 31773-95-2P

RL: ANT (Analyte); PUR (Purification or recovery); ANST (Analytical study); PREP (Preparation)

(antioxidant/anticancer potential of phenolic compds.

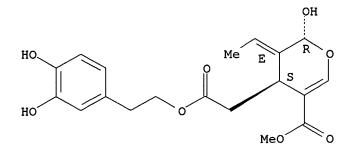
isolated from olive oil)

RN 31773-95-2 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-3,4-dihydro-2-hydroxy-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2R,3E,4S)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.



REFERENCE COUNT:

38 THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 9 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:319686 CAPLUS

DOCUMENT NUMBER: 132:339119

TITLE: Polyphenols: simple structures with high potential

AUTHOR(S): Metz, Gunter

CORPORATE SOURCE: Blaubeuren, 89143, Germany

SOURCE: Pharmazeutische Zeitung (2000), 145(16),

1273-1275,1278

CODEN: PHZIAP; ISSN: 0031-7136
PUBLISHER: Govi-Verlag Pharmazeutischer Verlag

DOCUMENT TYPE: Journal; General Review

LANGUAGE: German

AB A review with 6 refs. is given on the medicinal effects of polyphenols (e.g. anticarcinogen, antioxidative) including phenolic acids, cumarins and furocumarins, propolis, ingredients in olive oil, and ACA.

IT 32619-42-4, Oleuropein

RL: BAC (Biological activity or effector, except adverse); BOC (Biological occurrence); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); USES (Uses)

(medicinal effects of polyphenols)

RN 32619-42-4 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester,
(2S,3E,4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

REFERENCE COUNT:

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

CAPLUS COPYRIGHT 2006 ACS on STN ANSWER 10 OF 13

6

ACCESSION NUMBER: 1999:467227 CAPLUS

DOCUMENT NUMBER: 131:226102

TITLE: Studies on constituents with cytotoxic activity

from the stem bark of Syringa velutina

Park, Hee-Juhn; Lee, Myung-Sun; Lee, Kyung-Tae; AUTHOR(S):

Sohn, Il-Cheol; Han, Yong-Nam; Miyamoto, Ken-Ichi

Department of Botanical Resources, Sangi CORPORATE SOURCE:

University, Wonju, 220-702, S. Korea

Chemical & Pharmaceutical Bulletin (1999), 47(7), SOURCE:

1029-1031

CODEN: CPBTAL; ISSN: 0009-2363

Pharmaceutical Society of Japan PUBLISHER:

DOCUMENT TYPE: Journal English LANGUAGE:

Cytotoxic compds., oleuropein and a phenylethanoid glycoside (I) were isolated from the stem bark of Syringa velutina KOM. along with

coniferylaldehyde 4-0-glucoside, syringin, ligstroside,

(+)-syringaresinol 4-0-glucoside, (+)-medioresinol 4''-0-glucoside and

(-)-olivil 4''-O-glucoside. I was identified to be

3,4-dihydroxyphenylethyl alc. $8-O-\beta-D$ -glucopyranoside. Alc. $8-O-\beta-D$ -glucopyranoside. This compound showed the most potent cytotoxic effect on several tumor cell lines (P-388, L-1210,

SNU-5 and HL-60) among eight compds. isolated in the present study. We suggest that the 3,4-dihydroxyphenylethoxy moiety of this compound contributes to cytotoxicity.

IT 32619-42-4, Oleuropein

RL: BAC (Biological activity or effector, except adverse); BOC

Searcher Shears 571-272-2528 :

(Biological occurrence); BSU (Biological study, unclassified); BIOL (Biological study); OCCU (Occurrence)

(cytotoxic activities of constituents from stem bark of Syringa velutina)

RN 32619-42-4 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S,3E,4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

REFERENCE COUNT: 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 11 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1998:380086 CAPLUS

DOCUMENT NUMBER: 129:81138

TITLE: Free radical-scavenging properties of olive oil

polyphenols

AUTHOR(S): Visioli, Francesco; Bellomo, Giorgio; Galli,

Claudio

CORPORATE SOURCE: Institute of Pharmacological Sciences, University

of Milan, Italy

SOURCE: Biochemical and Biophysical Research

Communications (1998), 247(1), 60-64

CODEN: BBRCA9; ISSN: 0006-291X

PUBLISHER: Academic Press

DOCUMENT TYPE: Journal LANGUAGE: English

AB Plants in the Mediterranean basin, such as vine and olive trees, have developed an array of antioxidant defences to protect themselves from

environmental stress. Accordingly, the incidence of coronary heart disease and certain cancers is lower in the Mediterranean area, where olive oil is the dietary fat of choice. As opposed to other vegetable oils, extra virgin olive oil, which is obtained by phys. pressure from a whole fruit, is rich in phenolic components that are responsible for the particular stability of the oil. We have investigated the scavenging actions of some olive oil phenolics, namely hydroxytyrosol and oleuropein, with respect to superoxide anion generation, neutrophils respiratory burst, and hypochlorous acid. The low EC50s indicate that both compds. are potent scavengers of superoxide radicals and inhibitors of neutrophils respiratory burst: whenever demonstrated in vivo, these properties may partially explain the observed lower incidence of CHD and cancer associated with the Mediterranean diet.

IT 32619-42-4, Oleuropein

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study) (free radical-scavenging properties of olive oil polyphenols)

RN 32619-42-4 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S,3E,4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

REFERENCE COUNT:

37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 12 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1998:20210 CAPLUS

DOCUMENT NUMBER: 128:162831

TITLE: Oleuropein, the bitter principle of olives,

enhances nitric oxide production by mouse

macrophages

AUTHOR(S): Visioli, Francesco; Bellosta, Stefano; Galli,

Claudio

CORPORATE SOURCE: Institute of Pharmacological Sciences, Milan,

20133, Italy

SOURCE: Life Sciences (1998), 62(6), 541-546

CODEN: LIFSAK; ISSN: 0024-3205

PUBLISHER: Elsevier Science Inc.

DOCUMENT TYPE: Journal LANGUAGE: English

The Mediterranean diet, rich in fresh fruits and vegetables, has been associated with a lower incidence of cardiovascular disease and cancer, partly because of its high proportion of bioactive compds. such as vitamins, flavonoids and polyphenols. The major lipid component of such diet is the drupe-derived olive oil, that can be distinguished from other seed oils for the peculiar composition of its non-triglyceride fraction. In fact, several minor components, including polyphenols, grant the oil its particular taste and aroma. Oleuropein, the most abundant among these components, has been shown to be a potent antioxidant endowed with antiinflammatory properties. We investigated the effects of oleuropein on NO release in cell culture and its activity toward nitric oxide synthase (iNOS) expression. The results show that oleuropein dose-dependently enhance nitrite production in LPS-challenged mouse macrophages. This effect was blocked by the iNOS inhibitor L-NAME, indicating increased iNOS activity. Also, Western blot anal. of cell homogenates show that oleuropein increases iNOS expression in such cells. Taken together, our data suggest that, during endotoxin challenge, oleuropein potentiates the macrophage-mediated response, resulting in higher NO production, currently believed to be beneficial for cellular and organismal protection.

IT 32619-42-4, Oleuropein

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(oleuropein from olive oil enhances nitric oxide production by macrophages)

RN 32619-42-4 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S,3E,4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

REFERENCE COUNT:

THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 13 OF 13 CAPLUS COPYRIGHT 2006 ACS on STN

20

ACCESSION NUMBER:

1995:519455 CAPLUS

DOCUMENT NUMBER:

122:298024

TITLE:

Waste waters from olive oil production are rich in

natural antioxidants

AUTHOR(S):

Visioli, F.; Vinceri, F. F.; Galli, C.

CORPORATE SOURCE:

Inst. Pharmacological Sci., Univ. Milan, Milan,

I-20133, Italy

English

SOURCE:

LANGUAGE:

Experientia (1995), 51(1), 32-4 CODEN: EXPEAM; ISSN: 0014-4754

PUBLISHER: Birkhaeuser DOCUMENT TYPE: Journal

AB Milling of olive paste during olive oil production is accompanied by continuous washing with water, i.e. malaxation. The resulting wastewater is currently discarded. Since olives and olive oil are rich in natural antioxidants, we hypothesized that some of these might be extracted from the olive paste during malaxation. Interest in natural antioxidants is increasing because of the growing body of evidence indicating the involvement of oxygen-derived free radicals in several pathol. processes, such as cancer and atherosclerosis. A wastewater extract was characterized by HPLC and tested in a model of lipid peroxidn., copper sulfate-induced oxidation of low d. lipoproteins. The results demonstrate that wastewater exts. have powerful antioxidant activity and might therefore represent a cheap, as yet unused, source of antioxidants.

IT 32619-42-4, Oleuropein

RL: MSC (Miscellaneous)

(olive oil manufacturing wastewater antioxidants)

RN 32619-42-4 CAPLUS

CN 2H-Pyran-4-acetic acid, 3-ethylidene-2-(β-D-glucopyranosyloxy)-3,4-dihydro-5-(methoxycarbonyl)-, 2-(3,4-dihydroxyphenyl)ethyl ester, (2S,3E,4S)- (9CI) (CA INDEX NAME)

Absolute stereochemistry. Rotation (-). Double bond geometry as shown.

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L8 2 L4

L8 ANSWER 1 OF 2 CAOLD COPYRIGHT 2006 ACS on STN

AN CA64:8124e CAOLD

TI oleuropein - (III)

Panizzi, Luigi; Scarpati, M. L.; Trogolo, C.

ΑIJ

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4809-64-7 32619-42-4
IT
L8
    ANSWER 2 OF 2 CAOLD COPYRIGHT 2006 ACS on STN
    CA57:3398c CAOLD
AN
    constitution of oleuropein, a bitter glucoside of the olive with
TТ
    hypotensive action - (II)
    Panizzi, Luigi; Scarpati, M. L.; Oriente, G.
ΑU
    4751-57-9 7417-21-2 10597-60-1 32619-42-4 51856-81-6
IT
    90613-43-7. 91143-92-9 91556-07-9 91966-89-1 92038-78-3 92157-73-8
    92371-95-4 92730-99-9 92788-48-2 93404-86-5 96536-15-1 96650-87-2
FILE 'USPATFULL' ENTERED AT 12:34:48 ON 03 MAR 2006
CA INDEXING COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)
FILE COVERS 1971 TO PATENT PUBLICATION DATE: 2 Mar 2006 (20060302/PD)
FILE LAST UPDATED: 2 Mar 2006 (20060302/ED)
HIGHEST GRANTED PATENT NUMBER: US7007305
HIGHEST APPLICATION PUBLICATION NUMBER: US2006048257
CA INDEXING IS CURRENT THROUGH 28 Feb 2006 (20060228/UPCA)
ISSUE CLASS FIELDS (/INCL) CURRENT THROUGH: 2 Mar 2006 (20060302/PD)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Dec 2005
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Dec 2005
L9
            74 SEA ABB=ON PLU=ON L4
            27 SEA ABB=ON PLU=ON L9 AND (?CANCER? OR ?CARCIN? OR
L10
               ?TUMOUR? OR ?TUMOR? OR ?NEOPLAS?)
             3 SEA ABB=ON PLU=ON L10 NOT (PY=>2002 OR PD=>20021209)
L11
L11 ANSWER 1 OF 3 USPATFULL on STN
ACCESSION NUMBER:
                      2001:32806 USPATFULL
                      Water-soluble extract from olives
INVENTOR(S):
                      Crea, Roberto, San Mateo, CA, United States
                      Caglioti, Luciano, Rome, Italy
PATENT ASSIGNEE(S):
                      CreAgri L.L.C., Hayward, CA, United States (U.S.
                       corporation)
                          NUMBER KIND DATE
                       ______
                      US 6197308 B1 20010306
US 1999-359150 19990722
PATENT INFORMATION:
                                        19990722 (9)
APPLICATION INFO.:
                            NUMBER DATE
                       _____
PRIORITY INFORMATION: US 1998-93818P 19980723 (60)
DOCUMENT TYPE:
                      Utility
FILE SEGMENT:
                      Granted
PRIMARY EXAMINER:
                     Lilling, Herbert J.
LEGAL REPRESENTATIVE: Dehlinger, Peter J.
NUMBER OF CLAIMS:
EXEMPLARY CLAIM:
                      1
LINE COUNT:
                      309
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      The invention provides olive-derived vegetation water substantially
      free of monophenolic compounds (e.g., tyrosol and its derivatives)
      from olive pits. According to one aspect of the invention, the pits
      or seeds are removed from the olives prior to pressing. The pitless
      pulp or meat is then pressed to obtain a liquid-phase mixture
      including olive oil, vegetation water, and solid by-products. The
```

vegetation water is separated from the rest of the liquid-phase mixture and collected. The vegetation water is useful as a source of oleuropein.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L11 ANSWER 2 OF 3 USPATFULL on STN

ACCESSION NUMBER: 2000:174104 USPATFULL

Water-soluble extract from olives TITLE:

INVENTOR(S): Crea, Roberto, San Mateo, CA, United States

Caglioti, Luciano, Rome, Italy

CreAgri, Inc., Hayward, CA, United States (U.S. PATENT ASSIGNEE(S):

corporation)

NUMBER KIND DATE ______ PATENT INFORMATION: US 6165475 20001226 US 2000-491680 20000126 (9)

Continuation-in-part of Ser. No. US 1999-359150, RELATED APPLN. INFO.:

filed on 22 Jul 1999

NUMBER DATE

US 1998-93818P 19980723 (60) PRIORITY INFORMATION:

DOCUMENT TYPE: Utility FILE SEGMENT: Granted

FILE SEGMENT: Granted
PRIMARY EXAMINER: Lilling, Herbert J. LEGAL REPRESENTATIVE: Dehlinger, Peter J.

NUMBER OF CLAIMS: 7
EXEMPLARY CLAIM: 1
345

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

The invention provides a stable, olive-derived vegetation water substantially free of monophenolic compounds (e.g., tyrosol and its derivatives). The olive or seeds are removed from the olives prior to pressing. The pitless pulp or meat is then pressed to obtain a liquid-phase mixture including olive oil, vegetation water, and solid by-products. The vegetation water is separated from the rest of the liquid-phase mixture and collected, and stabilized for long term storage by acidification.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L11 ANSWER 3 OF 3 USPATFULL on STN

ACCESSION NUMBER: 2000:153273 USPATFULL

Use of an extract from the leaves of Olea Europea TITLE:

as an antiradical

INVENTOR(S): Amari, Giorgio, Milan, Italy

PATENT ASSIGNEE(S): B & T S.r.l., Milan, Italy (non-U.S. corporation)

NUMBER KIND DATE _____ PATENT INFORMATION: US 6146637 US 1999-250163 20001114 19990216 (9) APPLICATION INFO.:

NUMBER DATE _____ PRIORITY INFORMATION: IT 1998-MI317 19980219

DOCUMENT TYPE: Utility

FILE SEGMENT: Granted

PRIMARY EXAMINER: Lilling, Herbert J. LEGAL REPRESENTATIVE: Young & Thompson

NUMBER OF CLAIMS: 2
EXEMPLARY CLAIM: 1
LINE COUNT: 152

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The use of an extract from the leaves of Olea Europea as an antiradical. Preferably, the extract is added to a preparation wherein the concentration of the extract does not exceed 0.5% by weight. The extract can be used both for preparing cosmetic products, such as cosmetic creams—particularly sun protection creams—and for preparing alimentary products—particularly dietetic products.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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L12 406 SEA ABB=ON PLU=ON L4

L13 50 SEA ABB=ON PLU=ON L12 AND (CANCER? OR CARCIN? OR TUMOUR? OR TUMOR? OR NEOPLAS? OR ANTICANCER? OR ANTICARCIN? OR

ANTITUMOUR? OR ANTITUMOR? OR ANTINEOPLAS?)

L14 35 DUP REM L13 (15 DUPLICATES REMOVED)

L14 ANSWER 1 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on

STN

ACCESSION NUMBER: 2005:329527 BIOSIS DOCUMENT NUMBER: PREV200510114940

TITLE: Quantitation of oleuropein and related metabolites in

decoctions of Olea europaea leaves from ten Greek

cultivated varieties by HPLC with diode array detection

(HPLC-DAD).

AUTHOR(S): Agalias, Apostolis; Melliou, Eleni; Magiatis,

Prokopios; Mitaku, Sofia [Reprint Author]; Gikas,

Evagelos; Tsarbopoulos, Anthony

CORPORATE SOURCE: Univ Athens, Dept Pharm, Div Pharmacognosy and Nat Prod

Chem, Panepistimiopolis Zografou, GR-15771 Athens,

Greece

mitakou@pharm.uoa.gr

SOURCE: Journal of Liquid Chromatography & Related

Technologies, (2005) Vol. 28, No. 10, pp. 1557-1571.

ISSN: 1082-6076.

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 25 Aug 2005

Last Updated on STN: 25 Aug 2005

AB An extraction procedure and chromatographic methodology for the simultaneous quantitation of four major constituents in the boiling water extracts (decoctions) of Olea europaea leaves has been developed. The four studied constituents were oleuropein, elenolic acid, hydroxytyrosol, and tyrosol. The quantitation was performed using HPLC-DAD, whereas qualitative data were acquired using LC-MS.

The developed methodology was applied in the study-of ten Olea europaea varieties commonly cultivated in Greece. The chromatographic analysis revealed important differences among the varieties. The decoction of variety gaidouroelia was identified as the best source of oleuropein, but it was completely lacking of elenolic acid. The decoction of variety koronaiiki was the best source of hydroxytyrosol, whereas the variety mastoides was the best source of tyrosol and elenolic acid. In addition, the methanol and acetone extracts of one of the studied varieties (koranaiiki) were investigated, in order to compare the concentration of oleuropein in the extracts and the decoction. Interestingly, only a very low percent of the total oleuropein is present in the traditionally prepared decoction, while elenolic acid, which is a minor constituent of the extracts, was found to be one of the major constituents of the decoction.

L14 ANSWER 2 OF 35 EMBASE COPYRIGHT (c) 2006 Elsevier B.V. All rights reserved on STN

ACCESSION NUMBER: 2005486238 EMBASE

TITLE: Astonishing diversity of natural surfactants: 5.

Biologically active glycosides of aromatic metabolites.

AUTHOR: Dembitsky V.M.

CORPORATE SOURCE: V.M. Dembitsky, Department of Organic Chemistry, Hebrew

University, P.O. Box 39231, Jerusalem 91391, Israel.

dvalery@cc.huji.ac.il

SOURCE: Lipids, (2005) Vol. 40, No. 9, pp. 869-900. .

Refs: 328

ISSN: 0024-4201 CODEN: LPDSAP

COUNTRY: United States

DOCUMENT TYPE: Journal; General Review FILE SEGMENT: 030 Pharmacology

037 Drug Literature Index

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 20051128

Last Updated on STN: 20051128

AB This review article presents 342 aromatic glycosides, isolated from and identified in plants and microorganisms, that demonstrate different biological activities. They are of great interest, especially for the medicinal and/or pharmaceutical industries. These biologically active natural surfactants are good prospects for the future chemical preparation of compounds useful as antioxidant, anticancer, antimicrobial, and antibacterial agents. These glycosidic compounds have been classified into several groups, including simple aromatic compounds, stilbenes, phenylethanoids, phenylpropanoids, naphthalene derivatives, and anthracene derivatives. Copyright .COPYRGT. 2005 by AOCS Press.

L14 ANSWER 3 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN DUPLICATE 1

ACCESSION NUMBER: 2005:465993 BIOSIS DOCUMENT NUMBER: PREV200510246592

TITLE: Oleuropein, a non-toxic olive iridoid, is an anti-

tumor agent and cytoskeleton disruptor.

AUTHOR(S): Hamdi, Hamdi K. [Reprint Author]; Castellon, Raquel CORPORATE SOURCE: H2RC Corp, 1920 E Katella Ave, Suite U, Orange, CA 92867

USA

hkhamdi@mail.com

SOURCE: Biochemical and Biophysical Research Communications,

(SEP 2 2005) Vol. 334, No. 3, pp. 769-778.

CODEN: BBRCA9. ISSN: 0006-291X.

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 9 Nov 2005

Last Updated on STN: 9 Nov 2005

Oleuropein, a non-toxic secoiridoid derived from the olive tree, is a AB powerful antioxidant and anti-angiogenic agent. Here, we show it to be a potent anti-cancer compound, directly disrupting actin filaments in cells and in a cell-free assay. Oleuropein inhibited the proliferation and migration of advanced-grade human tumor cell lines in a dose-responsive manner. In a novel tube-disruption assay. Oleuropein irreversibly rounded cancer cells, preventing their replication, motility, and invasiveness; these effects were reversible in normal cells. When administered orally to mice that developed spontaneous tumors, Oleuropein completely regressed tumors in 9-12 days. When tumors were resected prior to complete regression, they lacked cohesiveness and had a crumbly consistency. No viable cells could be recovered from these tumors. These observations elevate Oleuropein from a non-toxic antioxidant into a potent antitumor agent with direct effects against tumor cells. Our data may also explain the cancer-protective effects of the olive-rich Mediterranean diet. (c) 2005 Elsevier Inc. All rights reserved.

L14 ANSWER 4 OF 35 EMBASE COPYRIGHT (c) 2006 Elsevier B.V. All rights reserved on STN

2005131586 EMBASE ACCESSION NUMBER:

Differential anti-inflammatory effects of phenolic TITLE:

compounds from extra virgin olive oil identified in

human whole blood cultures.

Miles E.A.; Zoubouli P.; Calder P.C. AUTHOR:

Dr. E.A. Miles, Institute of Human Nutrition, School of CORPORATE SOURCE:

Medicine, University of Southampton, Southampton,

United Kingdom. eam@soton.ac.uk.

SOURCE: Nutrition, (2005) Vol. 21, No. 3, pp. 389-394. .

Refs: 28

ISSN: 0899-9007 CODEN: NUTRER

COUNTRY: United States DOCUMENT TYPE: Journal; Article

Immunology, Serology and Transplantation FILE SEGMENT: 026

> 029 Clinical Biochemistry

English LANGUAGE: SUMMARY LANGUAGE: English

Entered STN: 20050407 ENTRY DATE:

Last Updated on STN: 20050407

Objective: The olive oil-rich Mediterranean diet protects against AB cardiovascular disease, which involves inflammatory processes. This study investigated the effects of phenolic compounds found in extra virgin olive oil on inflammatory mediator production by human mononuclear cells. Methods: Diluted human blood cultures were stimulated with lipopolysaccharide in the presence of phenolics (vanillic, p-coumaric, syringic, homovanillic and caffeic acids, kaempferol, oleuropein glycoside, and tyrosol) at concentrations of 10 (-7) to 10(-4) M. Concentrations of the inflammatory cytokines tumor necrosis factor- α , interleukin- 1β , and interleukin-6 and of the inflammatory eicosanoid prostaglandin E(2) were measured by enzyme-linked immunosorbent assay. Results: Oleuropein glycoside and caffeic acid decreased the concentration of

interleukin-1 β . At a concentration of 10(-4) M, oleuropein glycoside inhibited interleukin-1ß production by 80%, whereas caffeic acid inhibited production by 40%. Kaempferol decreased the concentration of prostaglandin E(2). At a concentration of 10 (-4) M, kaempferol inhibited prostaglandin E(2) production by 95%. No effects were seen on concentrations of interleukin-6 or tumor necrosis factor- α and there were no effects of the other phenolic compounds. Conclusions: Some, but not all, phenolic compounds derived from extra virgin olive oil decrease inflammatory mediator production by human whole blood cultures. This may contribute to the antiatherogenic properties ascribed to extra virgin olive oil. .COPYRGT. 2005 Elsevier Inc. All rights reserved.

L14 ANSWER 5 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN

DUPLICATE 2

ACCESSION NUMBER: 2005:330331 BIOSIS DOCUMENT NUMBER: PREV200510116584

The phenolic compounds of olive oil: structure,

biological activity and beneficial effects on human

Tripoli, Elisa; Giammanco, Marco [Reprint Author]; AUTHOR(S):

Tabacchi, Garden; Di Majo, Danila; Giammanco, Santo; La

Guardia, Maurizio

CORPORATE SOURCE: Univ Palermo, Fac Pharm, Inst Physiol and Human Nutr,

Via Augusto Elia 3, I-90127 Palermo, Italy

giammanco@unipa.it

Nutrition Research Reviews, (JUN 2005) Vol. 18, No. 1, SOURCE:

pp. 98-112.

CODEN: NREREX. ISSN: 0954-4224.

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 25 Aug 2005

Last Updated on STN: 25 Aug 2005

The Mediterranean diet is rich in vegetables, cereals, fruit, fish, AΒ milk, wine and olive oil and has salutary biological functions. Epidemiological studies have shown a lower incidence of atherosclerosis, cardiovascular diseases and certain kinds of cancer in the Mediterranean area. Olive oil is the main source of fat, and the Mediterranean diet's healthy effects can in particular be attributed not only to the high relationship between unsaturated and saturated fatty acids in olive oil but also to the antioxidant property of its phenolic compounds. The main phenolic compounds, hydroxytyrosol and oleuropein, which give extra-virgin olive oil its bitter, pungent taste, have powerful antioxidant activity both in vivo and in vitro. The present review focuses on recent works analysing the relationship between the structure of olive oil polyphenolic compounds and their antioxidant activity. These compounds' possible beneficial effects are due to their antioxidant activity, which is related to the development of atherosclerosis and cancer, and to anti-inflammatory and antimicrobial activity.

L14 ANSWER 6 OF 35 EMBASE COPYRIGHT (c) 2006 Elsevier B.V. All rights reserved on STN

ACCESSION NUMBER: 2005079028 EMBASE

The antioxidant properties of Greek foods and the TITLE:

flavonoid content of the Mediterranean menu.

AUTHOR: Vasilopoulou E.; Georga K.; Joergensen M.B.; Naska A.;

Trichopoulou A.

CORPORATE SOURCE: A. Trichopoulou, Department of Hygiene/Epidemiology,

> Shears 571-272-2528 Searcher :

School of Medicine, Natl./Kapodistrian Univ. of Athens,

Mikras Asias 75, Athens 115 27, Greece.

antonia@nut.uoa.gr

SOURCE: Current Medicinal Chemistry: Immunology, Endocrine and

Metabolic Agents, (2005) Vol. 5, No. 1, pp. 33-45. .

Refs: 117

ISSN: 1568-0134 CODEN: CMCIC8

COUNTRY: Netherlands

DOCUMENT TYPE: Journal; General Review

FILE SEGMENT: 016 Cancer

017 Public Health, Social Medicine and Epidemiology

018 Cardiovascular Diseases and Cardiovascular

Surgery

029 Clinical Biochemistry

030 Pharmacology

037 Drug Literature Index

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 20050303

Last Updated on STN: 20050303

The Mediterranean diet is currently attracting interest because of its AB health benefits that may be due, in part, to the high content of this diet in antioxidant phytochemicals. The variety and amount of phytochemicals taken with the consumption of primary and composite foods of the Mediterranean diet may provide better antiatherogenic properties than single phytochemicals. Flavonoids are the most important group of plant antioxidants. The Mediterranean diet is characterized by high intake of olive oil, fruit, vegetables, cereals, and legumes, some of which are good sources of flavonoids. Flavonoids consist of six principal classes: flavones, flavonols, flavan-3-ols, flavanones, anthocyanidins and isoflavones. The flavonoid intake from a traditional Greek plant-based weekly menu was calculated and the daily average flavonoid intake was found 118.6 mg/d, of which flavanones contribute 32% (38.5 mg/d), catechins (the most important group of flavan-3-ols) contribute 28% (32.7 mg/d), flavonols 22% (26.4 mg/d), anthocyanidins 9% (11 mg/d), flavones 8% (8.7 mg/d) and isoflavones contribute 1% (1.3 mg/d). Herbs and spices, which are commonly used in the traditional Greek cuisine, although added in small quantities, significantly contribute to the flavonol and flavone intake due to frequent consumption. The Greek version of the Mediterranean diet with its high consumption of fruit and vegetables is characterized by high intake of flavonoids in comparison to diets in northern European countries. . COPYRGT. 2005 Bentham Science Publishers Ltd.

L14 ANSWER 7 OF 35 MEDLINE on STN ACCESSION NUMBER: 2004559974 MEDLINE DOCUMENT NUMBER: PubMed ID: 15487893

TITLE: Acid-induced structural modifications of unsaturated

Fatty acids and phenolic olive oil constituents by

nitrite ions: a chemical assessment.

AUTHOR: Napolitano Alessandra; Panzella Lucia; Savarese Maria;

Sacchi Raffaele; Giudicianni Italo; Paolillo Livio;

d'Ischia Marco

CORPORATE SOURCE: Department of Organic Chemistry and Biochemistry,

University of Naples Federico II, Via Cinthia 4,

I-80126 Naples, Italy.. alesnapo@unina.it

SOURCE: Chemical research in toxicology, (2004 Oct) Vol. 17,

No. 10, pp. 1329-37.

Journal code: 8807448. ISSN: 0893-228X.

PUB. COUNTRY: United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 200503

ENTRY DATE: Entered STN: 20041110

Last Updated on STN: 20050330 Entered Medline: 20050329

AB The structural modifications of the unsaturated fatty acid components of triglycerides in extra virgin olive oil (EVOO) following exposure to nitrite ions in acidic media were determined by two-dimensional (2D) NMR spectroscopy, aided by (15)N labeling and GC analysis, allowing investigation of the matrix without fractionation steps. the presence of excess nitrite ions in a 1% sulfuric acid/oil biphasic system, extensive double bond isomerization of the oleic/linoleic acid components of triglycerides was observed associated with nitration/oxidation processes. Structurally modified species were identified as E/Z-nitroalkene, 1,2-nitrohydroxy, and 3-nitro-1-alkene(1,5-diene) derivatives based on (1)H, (13)C, and (15)N 2D NMR analysis in comparison with model compounds. Minor constituents of EVOO, including phenolic compounds and tocopherols, were also substantially modified by nitrite-derived nitrating species, even under milder reaction conditions relevant to those occurring in the gastric compartments. Novel nitrated derivatives of tyrosol, hydroxytyrosol, and oleuropein (6-8) were identified by LC/MS analysis of the polar fraction of EVOO and by comparison with synthetic samples. Overall, these results provide the first systematic description at the chemical level of the consequences of exposing EVOO to nitrite ions at acidic pH and offer an improved basis for further investigations in the field of toxic nitrosation/nitration reactions and dietary antinitrosating agents.

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ACCESSION NUMBER: 2005049849 EMBASE

TITLE: Olive oil and modulation of cell signaling in disease

prevention.

AUTHOR: Wahle K.W.J.; Caruso D.; Ochoa J.J.; Quiles J.L.

CORPORATE SOURCE: K.W.J. Wahle, School of Life Sciences, Robert Gordon

University, Aberdeen, AB25 1HG, United Kingdom.

k.wahle-1@rgu.ac.uk

SOURCE: Lipids, (2004) Vol. 39, No. 12, pp. 1223-1231. .

Refs: 86

ISSN: 0024-4201 CODEN: LPDSAP

COUNTRY: United States

DOCUMENT TYPE: Journal; Conference Article FILE SEGMENT: 029 Clinical Biochemistry

030 Pharmacology

037 Drug Literature Index

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 20050210

Last Updated on STN: 20050210

AB Epidemiological studies show that populations consuming a predominantly plant-based Mediterranean-style diet exhibit lower incidences of chronic diseases than those eating a northern European or North American diet. This observation has been attributed to the greater consumption of fruits and vegetables and the lower consumption

of animal products, particularly fat. Although total fat intake in Mediterranean populations can be higher than in other regions (ca. 40% of calories), the greater proportion is derived from olive oil and not animals. Increased olive oil consumption is implicated in a reduction in cardiovascular disease, rheumatoid arthritis, and, to a lesser extent, a variety of cancers. Olive oil intake also has been shown to modulate immune function, particularly the inflammatory processes associated with the immune system. Olive oil is a nonoxidative dietary component, and the attenuation of the inflammatory process it elicits could explain its beneficial effects on disease risk since oxidative and inflammatory stresses appear to be underlying factors in the etiology of these diseases in man. The antioxidant effects of olive oil are probably due to a combination of its high oleic acid content (low oxidation potential compared with linoleic acid) and its content of a variety of plant antioxidants, particularly oleuropein, hydroxytyrosol, and tyrosol. It is also possible that the high oleic acid content and a proportionate reduction in linoleic acid intake would allow a greater conversion of α-linolenic acid (18:3n-3) to longer-chain n-3 PUFA, which have characteristic health benefits. Adoption of a Mediterranean diet could confer health benefits in high-risk populations.

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DUPLICATE 3

ACCESSION NUMBER: 2004:114714 BIOSIS DOCUMENT NUMBER: PREV200400104817

TITLE: Herbal bioactivation: The good, the bad and the ugly. Zhou, Shufeng [Reprint Author]; Koh, Hwee-Ling; Gao, AUTHOR(S):

Yihuai; Gong, Zhi-yuan; Lee, Edmund Jon Deoon

Department of Pharmacy, Faculty of Science, National CORPORATE SOURCE:

University of Singapore, 18 Science Drive 4, Singapore,

117543, Singapore phazsf@nus.edu.sq

Life Sciences, (January 9 2004) Vol. 74, No. 8, pp. SOURCE:

935-968. print.

ISSN: 0024-3205 (ISSN print).

DOCUMENT TYPE: Article

General Review; (Literature Review)

LANGUAGE: English

Entered STN: 25 Feb 2004 ENTRY DATE:

Last Updated on STN: 25 Feb 2004

It has been well established that the formation of reactive AB metabolites of drugs is associated with drug toxicity. Similarly, there are accumulating data suggesting the role of the formation of reactive metabolites/intermediates through bioactivation in herbal toxicity and carcinogenicity. It has been hypothesized that the resultant reactive metabolites following herbal bioactivation covalently bind to cellular proteins and DNA, leading to toxicity via multiple mechanisms such as direct cytotoxicity, oncogene activation, and hypersensitivity reactions. This is exemplified by aristolochic acids present in Aristolochia spp, undergoing reduction of the nitro group by hepatic cytochrome P450 (CYP1A1/2) or peroxidases in extrahepatic tissues to reactive cyclic nitrenium ion. The latter was capable of reacting with DNA and proteins, resulting in activation of H-ras oncogene, gene mutation and finally carcinogenesis. Other examples are pulegone present in essential oils from many mint species; and teucrin A, a diterpenoid found in germander (Teuchrium chamaedrys) used as an adjuvant to slimming diets. Extensive pulegone metabolism generated p-cresol that was a glutathione depletory, and

> 571-272-2528 Searcher : Shears

the furan ring of the diterpenoids in germander was oxidized by CYP3A4 to reactive epoxide which reacts with proteins such as CYP3A and epoxide hydrolase. On the other hand, some herbal/dietary constituents were shown to form reactive intermediates capable of irreversibly inhibiting various CYPs. The resultant metabolites lead to CYP inactivation by chemical modification of the heme, the apoprotein, or both as a result of covalent binding of modified heme to the apoprotein. Some examples include bergamottin, a furanocoumarin of grapefruit juice; capsaicin from chili peppers; glabridin, an isoflavan from licorice root; isothiocyanates found in all cruciferous vegetables; oleuropein rich in olive oil; dially sulfone found in garlic; and resveratrol, a constituent of red wine. CYPs have been known to metabolize more than 95% therapeutic drugs and activate a number of procarcinogens as well. Therefore, mechanism-based inhibition of CYPs may provide an explanation for some reported herb-drug interactions and chemopreventive activity of herbs. Due to the wide use and easy availability of herbal medicines, there is increasing concern about herbal toxicity. The safety and quality of herbal medicine should be ensured through greater research, pharmacovigilance, greater regulatory control and better communication between patients and health professionals.

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ACCESSION NUMBER: 2004355686 EMBASE

TITLE: Olives and olive oil in cancer prevention.

AUTHOR: Owen R.W.; Haubner R.; Wurtele G.; Hull W.E.;

Spiegelhalder B.; Bartsch H.

CORPORATE SOURCE: R.W. Owen, Div. Toxicol. Cancer Risk Factors, German

Cancer Research Center, Im Neuenheimer Feld 280,

D-69120 Heidelberg, Germany. R.Owen@DKFZ-Heidelberg.de

SOURCE: European Journal of Cancer Prevention, (2004) Vol. 13,

No. 4, pp. 319-326. .

Refs: 28

ISSN: 0959-8278 CODEN: EJUPEK

COUNTRY:

United Kingdom
Journal; Article

DOCUMENT TYPE: FILE SEGMENT:

016 Cancer

030 Pharmacology

037 Drug Literature Index

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 20040909

Last Updated on STN: 20040909

Epidemiologic studies conducted in the latter part of the twentieth AB century demonstrate fairly conclusively that the people of the Mediterranean basin enjoy a healthy lifestyle with decreased incidence of degenerative diseases. The data show that populations within Europe that consume the so-called 'Mediterranean diet' have lower incidences of major illnesses such as cancer and cardiovascular disease. Studies have suggested that the health-conferring benefits of the Mediterranean diet are due mainly to a high consumption of fibre, fish, fruits and vegetables. More recent research has focused on other important factors such as olives and olive oil. Obviously fibre (especially wholegrain-derived products), fruits and vegetables supply an important source of dietary antioxidants. What is the contribution from olives and olive oil? Apparently the potential is extremely high but epidemiologic studies rarely investigate consumption of these very important products

in-depth, perhaps due to a lack of exact information on the types and amounts of antioxidants present. Recent studies have shown that olives and olive oil contain antioxidants in abundance. Olives (especially those that have not been subjected to the Spanish brining process) contain up to 16 g/kg typified by acteosides, hydroxytyrosol, tyrosol and phenyl propionic acids. Olive oil, especially extra virgin, contains smaller amounts of hydroxytyrosol and tyrosol, but also contains secoiridoids and lignans in abundance. Both olives and olive oil contain substantial amounts of other compounds deemed to be anticancer agents (e.g. squalene and terpenoids) as well as the peroxidation-resistant lipid oleic acid. It seems probable that olive and olive oil consumption in southern Europe represents an important contribution to the beneficial effects on health of the Mediterranean diet. .COPYRGT. 2004 Lippincott Williams & Wilkins.

L14 ANSWER 11 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN

ACCESSION NUMBER: 2004:350361 BIOSIS DOCUMENT NUMBER: PREV200400348082

TITLE: Production of highly purified hydroxytyrosol from Olea

europaea leaf extract biotransformed by

hyperthermophilic beta-glycosidase.

AUTHOR(S): Briante, Raffaella; Patumi, Maurizio; Febbraio,

Ferdinando; Nucci, Roberto [Reprint Author]

CORPORATE SOURCE: Ist Biochim Prot, CNR, Via Marconi 10, I-80125, Naples,

Italy

r.nucci@ibp.cnr.it

SOURCE: Journal of Biotechnology, (July 1 2004) Vol. 111, No.

1, pp. 67-77. print.

ISSN: 0168-1656 (ISSN print).

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 18 Aug 2004

Last Updated on STN: 18 Aug 2004

A large amount of highly purified hydroxytyrosol (91-94% in weight) is AB obtained in short time by a simple biotransformation of Olea europaea leaf extract by a partially purified hyperthermophilic beta-glycosidase immobilized on chitosan support. The biotransformation conditions have been modulated for increasing the hydroxytyrosol yield, whilst chitosan and chitin matrices are used as adsorbent materials in liquid phase hydroxytyrosol extraction from the biotransformed mixtures. Natural and non-toxic hydroxytyrosol has been by this way produced from a vegetal source, and this compound appeared for the first time highly purified by natural and biocompatible safe biopolymers in comparison to previous results. Moreover, the GC analyses have displayed that the eluates from a two-step bioreactor have qualitative composition very similar to that of the extra-virgin olive oil polar fraction. The proposed bioreactor could also find application in the utilization of olive mill waste waters (OMWW), medium rich in large amounts of oleuropein, which can be converted in pharmacologically active compounds. Copyright 2004 Elsevier B.V. All rights reserved.

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on STN

ACCESSION NUMBER: 2005:122403 BIOSIS DOCUMENT NUMBER: PREV200500125839

TITLE: Olive oil and oxidative stress.

AUTHOR(S): Visioli, Francesco [Reprint Author]; Bogani, Paola;

Grande, Simona; Gail, Claudio

CORPORATE SOURCE: Dept Pharmacol Sci, Univ Milan, Milan, Italy

francesco.visioli@unini.it

SOURCE: Grasas y Aceites, (January 2004) Vol. 55, No. 1, pp.

66-75. print.

ISSN: 0017-3495 (ISSN print).

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 1 Apr 2005

Last Updated on STN: 1 Apr 2005

In addition to the fatty acid profile of olive oil, which is high in the monounsaturated oleic acid and appears to be beneficial in reducing several risk factors for coronary heart disease and certain cancers, extra virgin olive oil contains a considerable amount of phenolic compounds, e.g. hydroxytyrosol and oleuropein, that are responsible for its peculiar taste and for its high stability. of evidence demonstrates that olive oil phenolics are powerful antioxidants. Although most of these studies have been carried out in vitro, some in vivo experiments confirm that olive oil phenolics are dose-dependently absorbed and that they retain their biological activities after ingestion. These data could in part explain the lower incidence of coronary heart disease in the Mediterranean area, where (extra virgin) olive oil is the principal source of fat.

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on STN

ACCESSION NUMBER: 2004:378026 BIOSIS PREV200400378017 DOCUMENT NUMBER:

Differential anti-inflammatory effects of phenolic TITLE:

compounds from olive oil identified in human whole

blood cultures.

Miles, E. A. [Reprint Author]; Zoubouli, R.; Calder, P. AUTHOR(S):

CORPORATE SOURCE: Sch MedInst Human Nutr, Univ Southampton, Southampton,

Hants, SO16 7PX, England

Chemistry and Physics of Lipids, (June 2004) Vol. 130, SOURCE:

No. 1, pp. 34-35. print.

Meeting Info.: 45th International Conference on the Bioscience of Lipids. Ioannina, Greece. May 25-29,

2004.

ISSN: 0009-3084 (ISSN print).

DOCUMENT TYPE:

Conference; (Meeting)

Conference; Abstract; (Meeting Abstract)

LANGUAGE:

English

ENTRY DATE:

Entered STN: 22 Sep 2004

Last Updated on STN: 22 Sep 2004

L14 ANSWER 14 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation DUPLICATE 4 on STN

ACCESSION NUMBER: 2004:368865 BIOSIS

DOCUMENT NUMBER: PREV200400370276

Involvement of oleuropein in (some) digestive metabolic TITLE:

pathways.

Polzonetti, V.; Egidi, D.; Vita, A.; Vincenzetti, S.; AUTHOR(S):

Natalini, P. [Reprint Author]

Dipartimento Sci Morfologiche and Biochim Comparate, CORPORATE SOURCE:

Univ Camerino, Via Camerini 2, I-62032, Camerino, Italy

paolo.natalini@unicam.it

SOURCE: Food Chemistry, (November 2004) Vol. 88, No. 1, pp.

> Shears 571-272-2528 Searcher

11-15. print.

CODEN: FOCHDJ. ISSN: 0308-8146.

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 8 Sep 2004

Last Updated on STN: 8 Sep 2004

Olive oil is the principal source of fats in the Mediterranean diet and it has been postulated that the components in olive oil can contribute to a lower incidence of coronary heart disease and cancers (prostate, colon, breast, and skin). The positive effects on human health can be attributed to the high level of phenolic compounds present in olive oil, the major ones being oleuropein, hydroxytyrosol and tyrosol. The aim of the present study was to evaluate the effect of oleuropein on enzymes involved in specific pathways of metabolism of proteins, carbohydrates and lipids. In particular, the effects of oleuropein on enzymes, such as trypsin, pepsin, lipase, glycerol dehydrogenase, glycerol-3-phosphate dehydrogenase, and glycerokinase, were investigated. Results demonstrate that oleuropein is able to activate pepsin and shows an inhibitory effect toward all the other enzymes tested, which suggests a new role for this polyphenol. In addition, a new method for lipase activity assay is presented. Copyright 2004 Elsevier Ltd, All rights reserved.

L14 ANSWER 15 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN DUPLICATE 5

ACCESSION NUMBER: 2003:497686 BIOSIS DOCUMENT NUMBER: PREV200300499640

TITLE: In vitro cytotoxicity to human cells in culture of some

phenolics from olive oil.

AUTHOR(S): Babich, H. [Reprint Author]; Visioli, F.

CORPORATE SOURCE: Department of Biology, Stern College for Women, Yeshiva

University, 245 Lexington Avenue, New York, NY, 10016,

USA

babich@ymail.yu.edu

SOURCE: Farmaco (Lausanne), (May 2003) Vol. 58, No. 5, pp.

403-407. print. ISSN: 0014-827X.

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 29 Oct 2003

Last Updated on STN: 29 Oct 2003

The neutral red in vitro cytotoxicity assay was used to evaluate the comparative responses of human cells isolated from tissues of the oral cavity to olive oil phenolics. The cell lines used included normal gingival fibroblasts, immortalized, nontumorigenic gingival epithelial cells, and carcinoma cells from the salivary gland. No differences in the relative sensitivities to the phenolics amongst the three cell types were noted. In general, for all cell types, the sequence of increasing cytotoxicity was: oleuropein aglycone>oleuropein glycoside, caffeic acid>o-coumaric acid>cinnamic acidmchgttyrosol, syringic acid, protocatechuic acid, vanillic acid. Cytotoxicity was noted only at phenolic concentrations far exceeding those attainable after habitual consumption, thus indicating that consumption of phenol-rich olive oil is safe.

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ACCESSION NUMBER: 2004280692 EMBASE

TITLE: Natural products and synthetic compounds as

immunomodulators.

AUTHOR: Kayser O.; Masihi K.N.; Kiderlen A.F.

CORPORATE SOURCE: Dr. A.F. Kiderlen, Robert Koch-Institut, Department of

Infectious Diseases, Cellular Defense Mechanisms Unit,

Nordufer 20, D-13353 Berlin, Germany

SOURCE: Expert Review of Anti-Infective Therapy, (2003) Vol. 1,

No. 2, pp. 319-335. .

Refs: 191

ISSN: 1478-7210 CODEN: ERATCK

COUNTRY: United Kingdom

DOCUMENT TYPE: Journal; General Review

FILE SEGMENT: 026 Immunology, Serology and Transplantation

029 Clinical Biochemistry

030 Pharmacology

037 Drug Literature Index

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 20040722

Last Updated on STN: 20040722

AB Research on immunomodulation by natural products or synthetic derivatives is of key interest for anti-infective therapy for a number of reasons. Many plant remedies well-known in traditional medicine or refined natural products in clinical use exert their anti-infective effects not only (if at all) by directly affecting the pathogen. At least part of their effect is indirect, by stimulating natural and adaptive defense mechanisms of the host. These findings have now given many empirical therapies a rationale, scientific basis and thereby a means for 'intelligent' improvement. In discovering the molecular mechanisms by which known remedies exert their effects, chosen elements further down the 'chain of command' might be synthesized and applied directly for more rapid and selective cure, omitting unwanted side effects. The direct use of recombinant cytokines, often in combination with antibiotics, is one consequence of this rationale. .COPYRGT. Future Drugs Ltd. All rights reserved.

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ACCESSION NUMBER: 2003069313 EMBASE

TITLE: Olive-oil consumption and cancer risk.

AUTHOR: Filik L.; Ozyilkan O.

SOURCE: European Journal of Clinical Nutrition, (1 Jan 2003)

Vol. 57, No. 1, pp. 191. .

Refs: 4

ISSN: 0954-3007 CODEN: EJCNEQ

COUNTRY: United Kingdom DOCUMENT TYPE: Journal; Letter FILE SEGMENT: 016 Cancer

017 Public Health, Social Medicine and Epidemiology

LANGUAGE: English

ENTRY DATE: Entered STN: 20030220

Last Updated on STN: 20030220

DATA NOT AVAILABLE FOR THIS ACCESSION NUMBER

L14 ANSWER 18 OF 35 MEDLINE on STN DUPLICATE 6

ACCESSION NUMBER: 2003067119 MEDLINE DOCUMENT NUMBER: PubMed ID: 12535851

TITLE: Simultaneous determination of oleuropein and

hydroxytyrosol in rat plasma using liquid

chromatography with fluorescence detection.

AUTHOR: Tan Hai-Wei; Tuck Kellie L; Stupans Ieva; Hayball Peter

J

CORPORATE SOURCE: Centre for Pharmaceutical Research, School of

Pharmaceutical, Molecular and Biomedical Sciences, University of South Australia, Adelaide, 5000,

Australia.

SOURCE: Journal of chromatography. B, Analytical technologies

in the biomedical and life sciences, (2003 Feb 25) Vol.

785, No. 1, pp. 187-91.

Journal code: 101139554. ISSN: 1570-0232.

PUB. COUNTRY: United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 200307

ENTRY DATE: Entered STN: 20030212

Last Updated on STN: 20030801 Entered Medline: 20030731

Oleuropein, the main glycoside present in olives, and hydroxytyrosol, AΒ the principal degradation product of oleuropein present in olive oil, have been linked to reduction of coronary heart disease and certain cancers. In the present study a direct and sensitive reversed-phase high-performance liquid chromatographic assay was developed for simultaneous quantification of both oleuropein and hydroxytyrosol. The plasma protein was precipitated with acetonitrile, samples were then centrifuged and supernatants were dried, and reconstituted with water prior to injection. The chromatographic analysis was carried out using a phenyl column and an isocratic elution of acidified water and acetonitrile with fluorescence detection at 281 and 316 nm for excitation and emission, respectively. The calibration curve was linear and limits of quantification were 30 ng/ml and 3 microg/ml for hydroxytyrosol and oleuropein, respectively. The method has been successfully applied to monitor oleuropein and hydroxytyrosol plasma levels in the rat.

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ACCESSION NUMBER: 2003243873 EMBASE TITLE: UV-induced skin damage.

AUTHOR: Ichihashi M.; Ueda M.; Budiyanto A.; Bito T.; Oka M.;

Fukunaga M.; Tsuru K.; Horikawa T.

CORPORATE SOURCE: M. Ichihashi, Division of Dermatology, Dept. of

Clinical Molecular Medicine, Kobe Univ. Grad. School of Medicine, 7-5-1, Kusunoki-cho, Chuo-ku, Kobe 650-0017,

Japan. ichihash@med.kobe.ac.jp

SOURCE: Toxicology, (15 Jul 2003) Vol. 189, No. 1-2, pp. 21-39.

Refs: 135

ISSN: 0300-483X CODEN: TXCYAC

COUNTRY: Ireland

DOCUMENT TYPE: Journal; General Review

FILE SEGMENT: 013 Dermatology and Venereology

016 Cancer

030 Pharmacology

037 Drug Literature Index

052 Toxicology

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 20030703

Last Updated on STN: 20030703

Solar radiation induces acute and chronic reactions in human and AR animal skin. Chronic repeated exposures are the primary cause of benign and malignant skin tumors, including malignant melanoma. Among types of solar radiation, ultraviolet B (290-320 nm) radiation is highly mutagenic and carcinogenic in animal experiments compared to ultraviolet A (320-400 nm) radiation. Epidemiological studies suggest that solar UV radiation is responsible for skin tumor development via gene mutations and immunosuppression, and possibly for photoaging. In this review, recent understanding of DNA damage caused by direct UV radiation and by indirect stress via reactive oxygen species (ROS) and DNA repair mechanisms, particularly nucleotide excision repair of human cells, are discussed. In addition, mutations induced by solar UV radiation in p53, ras and patched genes of non-melanoma skin cancer cells, and the role of ROS as both a promoter in UVcarcinogenesis and an inducer of UV-apoptosis, are described based primarily on the findings reported during the last decade. Furthermore, the effect of UV on immunological reaction in the skin is discussed. Finally, possible prevention of UV-induced skin cancer by feeding or topical use of antioxidants, such as polyphenols, vitamin C, and vitamin E, is discussed. .COPYRGT. 2003 Published by Elsevier Science Ireland Ltd.

L14 ANSWER 20 OF 35 EMBASE COPYRIGHT (c) 2006 Elsevier B.V. All rights reserved on STN

ACCESSION NUMBER: 2003035183 EMBASE

TITLE: The randomized controlled trial in studies using

biomarkers.

AUTHOR: Vineis P.

CORPORATE SOURCE: P. Vineis, Dipt. di Sci. Biomed. e Oncol. Umana,

University of Torino, via Santena 7, Torino, Italy.

paolo.vineis@unito.it

SOURCE: Biomarkers, (2003) Vol. 8, No. 1, pp. 13-32...

Refs: 21

ISSN: 1354-750X CODEN: BIOMFA

COUNTRY: United Kingdom

DOCUMENT TYPE: Journal; General Review

FILE SEGMENT: 016 Cancer

017 Public Health, Social Medicine and Epidemiology

028 Urology and Nephrology 029 Clinical Biochemistry

030 Pharmacology

037 Drug Literature Index

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 20030130

Last Updated on STN: 20030130

AB The randomized controlled trial (RCT) is a scientific experiment during which observations on the effects of therapy or a preventive action are conducted by the researcher under rigorous control. The purpose of the experiment is to clear the uncertainties surrounding a clinical/research issue and involves isolating the 'treatment' and 'end result' variables from external influences. RCTs therefore make use of scientific method standards: measuring, which includes the possibility of reproducing observations; controlling factors unconnected to the cause-effect relationship of interest; and the external verification or 'falsification' of the cause-effect

relationship. Many RCTs are now including biomarkers to answer scientific questions in a more accurate way. In the present methodological paper, the main aspects involved in the design and conduction of a trial are discussed, with special emphasis on the use of biomarkers. Aspects that are often overlooked by scientists involved in the design of trials include multiple comparisons, subgroup analysis, the duration of the observations, the use of surrogate endpoints, and ethical issues. This review summarizes the main issues that should be addressed in a protocol, and illustrates these with an example.

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ACCESSION NUMBER: 2002286207 EMBASE

TITLE: Exocyclic DNA adducts as oxidative stress markers in

colon carcinogenesis: Potential role of lipid peroxidation, dietary fat and antioxidants.

AUTHOR: Bartsch H.; Nair J.; Owen R.W.

CORPORATE SOURCE: H. Bartsch, Div. of Toxicol./Cancer Risk Factors,

German Cancer Research Center DKFZ, Im Neuenheimer Feld

280, D-69120 Heidelberg, Germany

SOURCE: Biological Chemistry, (2002) Vol. 383, No. 6, pp.

915-921. . Refs: 47

ISSN: 1431-6730 CODEN: BICHF3

COUNTRY: Germany

DOCUMENT TYPE: Journal; General Review

FILE SEGMENT: 005 General Pathology and Pathological Anatomy

016 Cancer

030 Pharmacology

037 Drug Literature Index

048 Gastroenterology

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 20020829

Last Updated on STN: 20020829

Molecular pathways to colorectal cancer involve multiple AB genetic changes, whereby extensive oxyradical damage causes mutations in cancer-related genes and leads to a cycle of cell death and regenaration. Besides direct oxidative DNA-damage, reactive oxygen and nitrogen species can induce etheno (E)-DNA adducts mainly via trans-4-hydroxy-2-nonenal, generated as the major aldehyde by lipid peroxidation (LPO) of $\omega-6$ PUFAs. Patients with familial adenomatous polyposis (FAP) develop multiple colorectal adenomas. In affected tissues increased LPO could be trigered due to increased arachidonic acid metabolism as a result of elevated cyclooxygenase. Our studies demonstrated an increased ε-DNA adduct level in affected colon epithelia of FAP patients. ε-DNA adducts are promutagenic and can cause genomic instability that drives colorectal adenoma to malignancy. We have further investigated the potential chemopreventive properties of olive oil and its polyphenolic components. 'Mediterranean diet', of which olive oil is a major fatty acid source, has protective effects against human breast and colorectal cancers. Olive oil extracts and the newly identified lignan fractions showed high antioxidant capacity in vitro. As ϵ -DNA adducts are biomarkers for oxidative stress and LPO induced DNA damage, they can verify the efficacy of newly identified antioxidants, e.g. from olive oil, as chemopreventive agents against colon carcinogenesis.

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DUPLICATE 7 on STN

2003:57519 BIOSIS ACCESSION NUMBER: PREV200300057519 DOCUMENT NUMBER:

Major phenolic compounds in olive oil: Metabolism and TITLE:

health effects.

Tuck, Kellie L. [Reprint Author]; Hayball, Peter J. AUTHOR(S):

CORPORATE SOURCE: Centre for Pharmaceutical Research, School of

Pharmaceutical, Molecular and Biomedical Sciences, University of South Australia, Adelaide, 5000,

Australia

kellie.tuck@unisa.edu.au

Journal of Nutritional Biochemistry, (November 2002) SOURCE:

Vol. 13, No. 11, pp. 636-644. print.

CODEN: JNBIEL. ISSN: 0955-2863.

DOCUMENT TYPE: Article

General Review; (Literature Review)

LANGUAGE: English

ENTRY DATE: Entered STN: 22 Jan 2003

Last Updated on STN: 22 Jan 2003

It has been postulated that the components in olive oil in the Mediterranean diet, a diet which is largely vegetarian in nature, can contribute to the lower incidence of coronary heart disease and prostate and colon cancers. The Mediterranean diet includes the consumption of large amounts of olive oil. Olive oil is a source of at least 30 phenolic compounds. The major phenolic compounds in olive oil are oleuropein, hydroxytyrosol and tyrosol. Recently there has been a surge in the number of publications that has investigated their biological properties. The phenolic compounds present in olive oil are strong antioxidants and radical scavengers. Olive "waste water" also possesses compounds which are strong antioxidant and radical scavengers. Typically, hydroxytyrosol is a superior antioxidant and radical scavenger to oleuropein and tyrosol. Hydroxytyrosol and oleuropein have antimicrobial activity against ATTC bacterial strains and clinical bacterial strains. Recent syntheses of labeled and unlabelled hydroxytyrosol coupled with superior analytical techniques have enabled its absorption and metabolism to be studied. It has recently been found that hydroxytyosol is renally excreted unchanged and as the following metabolites as its glucuronide conjugate, sulfate conjugate, homovanillic acid, homovanillic alcohol, 3,4-dihydroxyphenylacetic acid and 3,4-dihydroxyphenylacetaldehyde. Studies with tyrosol have shown that it is excreted unchanged and as its conjugates. This review summarizes the antioxidant abilities; the scavenging abilities and the biological fates of hydroxytyrosol, oleuropein and tyrosol which have been published in recent years.

L14 ANSWER 23 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation DUPLICATE 8 on STN

2002:535154 BIOSIS ACCESSION NUMBER: DOCUMENT NUMBER: PREV200200535154

TITLE: Olive oil phenolics: Effects on DNA-oxidation and redox

enzyme mRNA in prostate cells.

AUTHOR(S): Lund, Elizabeth [Reprint author]

Institute of Food Research, Norwich Research Park, CORPORATE SOURCE:

Colney, Norwich, NR4 7UA, UK

British Journal of Nutrition, (September, 2002) Vol. SOURCE:

88, No. 3, pp. 223-224. print. CODEN: BJNUAV. ISSN: 0007-1145.

Shears 571-272-2528 Searcher :

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 16 Oct 2002

Last Updated on STN: 5 Dec 2002

L14 ANSWER 24 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation

on STN

ACCESSION NUMBER: 2002:421390 BIOSIS DOCUMENT NUMBER: PREV200200421390

TITLE: Biological properties of olive oil phytochemicals. AUTHOR(S): Visioli, Francesco [Reprint author]; Galli, Claudio University of Milan, Institute of Pharmacological CORPORATE SOURCE:

Sciences, Via Balzaretti 9, 20133, Milan, Italy

francesco.visioli@unimi.it

Critical Reviews in Food Science and Nutrition, (May, SOURCE:

2002) Vol. 42, No. 3, pp. 209-221. print.

ISSN: 1040-8398.

Article DOCUMENT TYPE:

General Review; (Literature Review)

LANGUAGE: English

Entered STN: 7 Aug 2002 ENTRY DATE:

Last Updated on STN: 7 Aug 2002

L14 ANSWER 25 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation

on STN

ACCESSION NUMBER: 2002:143530 BIOSIS PREV200200143530 DOCUMENT NUMBER:

Antioxidant and other biological activities of phenols TITLE:

from olives and olive oil.

Visioli, Francesco [Reprint author]; Poli, Andrea; AUTHOR(S):

Galli, Claudio

Department of Pharmacological Sciences, University of CORPORATE SOURCE:

Milan, Via Balzaretti 9, 20133, Milan, Italy

francesco.visioli@unimi.it

SOURCE: Medicinal Research Reviews, (January, 2002) Vol. 22,

No. 1, pp. 65-75. print.

CODEN: MRREDD. ISSN: 0198-6325.

DOCUMENT TYPE: Article LANGUAGE: English

Entered STN: 14 Feb 2002 ENTRY DATE:

Last Updated on STN: 26 Feb 2002

L14 ANSWER 26 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation

on STN

2001:417654 BIOSIS ACCESSION NUMBER: DOCUMENT NUMBER: PREV200100417654

Water-soluble extract from olives. TITLE:

Crea, Roberto [Inventor]; Caglioti, Luciano [Inventor, AUTHOR(S):

Reprint author]

CORPORATE SOURCE: Rome, Italy

ASSIGNEE: CreAgri L.L.C., Hayward, CA, USA

PATENT INFORMATION: US 6197308 20010306

Official Gazette of the United States Patent and SOURCE:

Trademark Office Patents, (Mar. 6, 2001) Vol. 1244, No.

1. e-file.

CODEN: OGUPE7. ISSN: 0098-1133.

DOCUMENT TYPE: Patent LANGUAGE: English

ENTRY DATE: Entered STN: 29 Aug 2001

Shears 571-272-2528 Searcher :

Last Updated on STN: 22 Feb 2002

The invention provides olive-derived vegetation water substantially AB free of monophenolic compounds (e.g., tyrosol and its derivatives) from olive pits. According to one aspect of the invention, the pits or seeds are removed from the olives prior to pressing. The pitless pulp or meat is then pressed to obtain a liquid-phase mixture including olive oil, vegetation water, and solid by-products. vegetation water is separated from the rest of the liquid-phase mixture and collected. The vegetation water is useful as a source of oleuropein.

L14 ANSWER 27 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation DUPLICATE 9 on STN

2001:522398 BIOSIS ACCESSION NUMBER: DOCUMENT NUMBER: PREV200100522398

The inhibitory effects of compounds from olive leaf on TITLE:

tumor necrosis factor production and on

beta-hexosaminidase release.

AUTHOR(S): Nishibe, Sansei [Reprint author]; Han, Yingmei [Reprint

author]; Noguchi, Yukari [Reprint author]; Ueda,

Hiroshi; Yamazaki, Masatoshi; Mizutani, Kenji; Kambara,

Toshimitsu; Kishida, Naoko

Faculty of Pharmaceutical Sciences, Health Sciences CORPORATE SOURCE:

University of Hokkaido, Ishikari-Tobetsu, Hokkaido,

061-0293, Japan

SOURCE: Natural Medicines, (August, 2001) Vol. 55, No. 4, pp.

> 205-208. print. ISSN: 1340-3443.

DOCUMENT TYPE: Article English LANGUAGE:

Entered STN: 7 Nov 2001 ENTRY DATE:

Last Updated on STN: 23 Feb 2002

The extraction and isolation of olive leaf gave luteolin AΒ

7-O-glucoside, luteolin 4'-O-glucoside and oleuropein as the major

components. The inhibitory effects of these compounds on

tumor necrosis factor (TNF-alpha) production and on

beta-hexosaminidase release from rat basophilic leukemia (RBL-2H3) cells, which were both recently found to be linked to allergic

reaction, were examined. Oleuropein showed a potent inhibitory effect on TNF-alpha production. Luteolin 4'-O-glucoside showed a strong inhibitory effect on beta-hexosaminidase release (IC50: 17.1 mug/ml).

L14 ANSWER 28 OF 35 MEDLINE on STN DUPLICATE 10

ACCESSION NUMBER: 2001098429 MEDLINE PubMed ID: 11110859 DOCUMENT NUMBER:

Oleuropein, an antioxidant polyphenol from olive oil, TITLE:

is poorly absorbed from isolated perfused rat

intestine.

Edgecombe S C; Stretch G L; Hayball P J AUTHOR:

CORPORATE SOURCE: Centre for Pharmaceutical Research, University of South

Australia, North Terrace, Adelaide, South Australia,

5000, Australia.

The Journal of nutrition, (2000 Dec) Vol. 130, No. 12, SOURCE:

pp. 2996-3002.

Journal code: 0404243. ISSN: 0022-3166.

PUB. COUNTRY:

United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

> Shears 571-272-2528 Searcher :

ENTRY MONTH: 200102

ENTRY DATE: Entered STN: 20010322

Last Updated on STN: 20010322 Entered Medline: 20010201

Epidemiological studies have shown that the incidence of heart disease AB and certain cancers is lower in the Mediterranean region. This has been attributed to the high consumption of olive oil in the Mediterranean diet, which contains polyphenolic compounds with antioxidant activity. Although many in vitro studies have been performed to elucidate mechanisms by which these compounds may act, there are virtually no data relating to their fate after ingestion. Therefore, we decided to investigate the intestinal absorption of one of the major olive oil polyphenolics, oleuropein. To do this, a novel in situ intestinal perfusion technique was developed, and the absorption of oleuropein was studied under both iso-osmotic and hypotonic luminal conditions. Oleuropein was absorbed, with an apparent permeability coefficient (P:(app)) of $1.47 +/-0.13 \times 10(-6)$ cm/s (+/-SE) observed under iso-osmotic conditions. The mechanism of absorption is unclear but may involve transcellular transport (SGLT1) or paracellular movement. Under hypotonic conditions, the permeability of oleuropein was significantly greater (5.92 \pm -0.49 x 10(-6) cm/s, P: < 0.001). This increase is thought to be due to an increase in paracellular movement facilitated by the opening of paracellular junctions in response to hypotonicity. Overall, we determined that the olive oil polyphenolic oleuropein can be absorbed, albeit poorly, from isolated perfused rat intestine. Therefore, it is possible that it or its metabolites may confer a positive health benefit after the consumption of olive oil, most likely via an antioxidant mechanism.

L14 ANSWER 29 OF 35 EMBASE COPYRIGHT (c) 2006 Elsevier B.V. All rights reserved on STN

ACCESSION NUMBER: 2000409319 EMBASE

TITLE: Protective effect of topically applied olive oil

against photocarcinogenesis following UVB exposure of

mice.

AUTHOR: Budiyanto A.; Ahmed N.U.; Wu A.; Bito T.; Nikaido O.;

Osawa T.; Ueda M.; Ichihashi M.

CORPORATE SOURCE: M. Ueda, Department of Dermatology, Kobe University

School of Medicine, 7-5-1 Kusunoki-cho, Chuo-ku, Kobe

650-0017, Japan. mueda@med.kobe-u.ac.jp

SOURCE: Carcinogenesis, (2000) Vol. 21, No. 11, pp. 2085-2090.

Refs: 31

ISSN: 0143-3334 CODEN: CRNGDP

COUNTRY: United Kingdom
DOCUMENT TYPE: Journal; Article

FILE SEGMENT: 013 Dermatology and Venereology

016 Cancer

030 Pharmacology

037 Drug Literature Index

052 Toxicology

LANGUAGE: English SUMMARY LANGUAGE: English

ENTRY DATE: Entered STN: 20001213

Last Updated on STN: 20001213

AB Reactive oxygen species have been shown to play a role in ultraviolet light (UV)-induced skin carcinogenesis. Vitamin E and green tea polyphenols reduce experimental skin cancers in mice

mainly because of their antioxidant properties. Since olive oil has also been reported to be a potent antioxidant, we examined its effect on UVB-induced skin carcinogenesis in hairless mice. Extra-virgin olive oil was applied topically before or after repeated exposure of mice to UVB. The onset of UVB-induced skin tumors was delayed in mice painted with olive oil compared with UVB control mice. However, with increasing numbers of UVB exposures, differences in the mean number of tumors between UVB control mice and mice pretreated with olive oil before UVB exposure (pre-UVB group) were lost. In contrast, mice that received olive oil after UVB exposure (post-UVB group) showed significantly lower numbers of tumors per mouse than those in the UVB control group throughout the experimental period. The mean number of tumors per mouse in the UVB control, pre-UVB and post-UVB groups was 7.33, 6.69 and 2.64, respectively, in the first experiment, and 8.53, 9.53 and 3.36 in the second experiment. Camellia oil was also applied, using the same experimental protocol, but did not have a suppressive effect. Immunohistochemical analysis of DNA damage in the form of cyclobutane pyrimidine dimers (CPD), (6-4) photoproducts and 8-hydroxy-2'-deoxyguanosine (8-OHdG) in samples taken 30 min after a single exposure of UVB showed no significant difference between UVB-irradiated control mice and the pre-UVB group. In the post-UVB group, there were lower levels of 8-OHdG in epidermal nuclei, but the formation of CPD and (6-4) photoproducts did not differ. Exposure of olive oil to UVB before application abrogated the protective effect on 8-OHdG formation. These results indicate that olive oil topically applied after UVB exposure can effectively reduce UVB-induced murine skin tumors, possibly via its anti-oxidant effects in reducing DNA damage by reactive oxygen species, and that the effective component may be labile to UVB.

L14 ANSWER 30 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN

ACCESSION NUMBER: 2000:236012 BIOSIS DOCUMENT NUMBER: PREV200000236012

TITLE: Skin anti-inflammatory activity of hydroxytyrosol and

its acid form 3,4 dihydroxyphenylacetic acid.

AUTHOR(S): Despotopoulos, A. [Reprint author]; Rallis, M. [Reprint

author]; Marakos, P. [Reprint author]; Rodis, P.;
Proxenia, N.; Demetzos, C. [Reprint author]; Xenos, K.;

Katsarou, A.; Tsaldaris, I. [Reprint author];

Papaioannou, G. [Reprint author]

CORPORATE SOURCE: University of Athens, Athens, Greece

SOURCE: Journal of Investigative Dermatology, (April, 2000)

Source: Sournal of investigative Definatorogy, (April, 2000)

Vol. 114, No. 4, pp. 881. print.

Meeting Info.: 61st Annual Meeting of the Society for Investigative Dermatology. Chicago, Illinois, USA. May

10-14, 2000.

CODEN: JIDEAE. ISSN: 0022-202X.

DOCUMENT TYPE: Conference; (Meeting)

Conference; Abstract; (Meeting Abstract)

LANGUAGE: English

ENTRY DATE: Entered STN: 7 Jun 2000

Last Updated on STN: 5 Jan 2002

L14 ANSWER 31 OF 35 EMBASE COPYRIGHT (c) 2006 Elsevier B.V. All rights reserved on STN

ACCESSION NUMBER: 2000163671 EMBASE

TITLE: [Polyphenols: Simple structures with high potency].

POLYPHENOLE: EINFACHE STRUKTUREN MIT HOHEM POTENZIAL.

AUTHOR: Metz G.

Dr. G. Metz, Auf dem Rucken 29, 89146 Blaubeuren, CORPORATE SOURCE:

Germany

Pharmazeutische Zeitung, (20 Apr 2000) Vol. 145, No. SOURCE:

16, pp. 23-28. .

Refs: 6

ISSN: 0031-7136 CODEN: PZSED5

COUNTRY:

Germany Journal; (Short Survey)

DOCUMENT TYPE: FILE SEGMENT:

030 Pharmacology

037 Drug Literature Index

LANGUAGE:

German

ENTRY DATE:

Entered STN: 20000525

Last Updated on STN: 20000525

DATA NOT AVAILABLE FOR THIS ACCESSION NUMBER

L14 ANSWER 32 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation

on STN

ACCESSION NUMBER: 2000:397299 BIOSIS PREV200000397299 DOCUMENT NUMBER:

Studies on constituents with cytotoxic activity from TITLE:

the stem bark of Syringa velutina.

Park, Hee-Juhn; Lee, Myung-Sun; Lee, Kyung-Tae; Sohn, AUTHOR(S):

Il-Cheol; Han, Yong-Nam; Miyamoto, Ken-ichi [Reprint

Department of Pharmacology and Pharmaceutics, Graduate CORPORATE SOURCE:

School of Natural Science and Technology, Kanazawa University, 13-1 Takara-machi, Kanazawa, 920-9042,

Chemical and Pharmaceutical Bulletin (Tokyo), (July, SOURCE:

1999) Vol. 47, No. 7, pp. 1029-1031. print.

CODEN: CPBTAL. ISSN: 0009-2363.

DOCUMENT TYPE:

Article

LANGUAGE: ENTRY DATE: English Entered STN: 20 Sep 2000

Last Updated on STN: 8 Jan 2002

Cytotoxic compounds, oleuropein (1) and a phenylethanoid glycoside (2) were isolated from the stem bark of Syringa velutina KOM. along with

coniferylaldehyde 4-O-glucoside, syringin, ligstroside,

(+)-syringaresinol 4-0-glucoside, (+)-medioresinol 4'-0-glucoside and (-)-olivil 4'-O-glucoside. Phenylethanoid glycoside (2) was identified to be 3,4-dihydroxyphenylethyl alcohol 8-O-beta-Dglucopyranoside. This compound showed the most potent cytotoxic

effect on several tumor cell lines (P-388, L-1210, SNU-5 and HL-60) among eight compounds isolated in the present study. We suggest that the 3,4-dihydroxyphenylethoxy moiety of this compound contributes to cytotoxicity.

MEDLINE on STN DUPLICATE 11 L14 ANSWER 33 OF 35

MEDLINE ACCESSION NUMBER: 1998124113 DOCUMENT NUMBER: PubMed ID: 9464466

TITLE: Oleuropein, the bitter principle of olives, enhances

nitric oxide production by mouse macrophages.

Visioli F; Bellosta S; Galli C AUTHOR:

Institute of Pharmacological Sciences, Milan, Italy.. CORPORATE SOURCE:

Francesco. Visioli@unimi.it

SOURCE: Life sciences, (1998) Vol. 62, No. 6, pp. 541-6.

Journal code: 0375521. ISSN: 0024-3205.

Shears 571-272-2528 Searcher :

PUB. COUNTRY: ENGLAND: United Kingdom

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 199802

ENTRY DATE: Entered STN: 19980226

Last Updated on STN: 19980226 Entered Medline: 19980219

The Mediterranean diet, rich in fresh fruits and vegetables, has been AB associated with a lower incidence of cardiovascular disease and cancer, partly because of its high proportion of bioactive compounds such as vitamins, flavonoids and polyphenols. The major lipid component of such diet is the drupe-derived olive oil, that can be distinguished from other seed oils for the peculiar composition of its non-triglyceride fraction. In fact, several minor components, including polyphenols, grant the oil its particular taste and aroma. Oleuropein, the most abundant among these components, has been shown to be a potent antioxidant endowed with antiinflammatory properties. We investigated the effects of oleuropein on NO release in cell culture and its activity toward nitric oxide synthase (iNOS) expression. The results show that oleuropein dose-dependently enhance nitrite production in LPS-challenged mouse macrophages. This effect was blocked by the iNOS inhibitor L-NAME, indicating increased iNOS activity. Also, Western blot analysis of cell homogenates show that oleuropein increases iNOS expression in such cells. Taken together, our data suggest that, during endotoxin challenge, oleuropein potentiates the macrophage-mediated response, resulting in higher NO production, currently believed to be beneficial for cellular and organismal protection.

L14 ANSWER 34 OF 35 MEDLINE on STN ACCESSION NUMBER: 1998456830 MEDLINE DOCUMENT NUMBER: PubMed ID: 9786644

TITLE: Cytostatic activity of some compounds from the

unsaponifiable fraction obtained from virgin olive oil.

AUTHOR: Saenz M T; Garcia M D; Ahumada M C; Ruiz V

CORPORATE SOURCE: Laboratori do Farmacognosia, Universidad de Sevilla,

41012 Seville, Spain.

SOURCE: Farmaco (Societa chimica italiana : 1989), (1998 Jun

30) Vol. 53, No. 6, pp. 448-9.

Journal code: 8912641. ISSN: 0014-827X.

PUB. COUNTRY: Italy

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 199810

ENTRY DATE: Entered STN: 19981029

Last Updated on STN: 19981029 Entered Medline: 19981022

AB Oleuropein, tyrosol, squalene and the fraction of sterols and triterpenoid dialcohols from the unsaponifiable fraction obtained from virgin olive oil have been tested for possible cytostatic activity against McCoy cells, using 6-mercaptopurine as a positive control. The samples of sterols and triterpenic dialcohols showed a strong activity.

L14 ANSWER 35 OF 35 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN DUPLICATE 12

ACCESSION NUMBER: 1998:323225 BIOSIS

DOCUMENT NUMBER: PREV199800323225

TITLE: Free radical-scavenging properties of olive oil

polyphenols.

AUTHOR(S): Visioli, Francesco [Reprint author]; Bellomo, Giorgio;

Galli, Claudio

CORPORATE SOURCE: Inst. Pharmacological Sci., Univ. Milan, Milan, Italy

SOURCE: Biochemical and Biophysical Research Communications,

(June 9, 1998) Vol. 247, No. 1, pp. 60-64. print.

CODEN: BBRCA9. ISSN: 0006-291X.

DOCUMENT TYPE: Article LANGUAGE: English

ENTRY DATE: Entered STN: 22 Jul 1998

Last Updated on STN: 22 Jul 1998

Plants in the Mediterranean basin, such as vine and olive trees, have AB developed an array of antioxidant defenses to protect themselves from environmental stress. Accordingly, the incidence of coronary heart disease and certain cancers is lower in the Mediterranean area, where olive oil is the dietary fat of choice. As opposed to other vegetable oils, extra virgin olive oil, which is obtained by physical pressure from a whole fruit, is rich in phenolic components that are responsible for the particular stability of the oil. We have investigated the scavenging actions of some olive oil phenolics, namely hydroxytyrosol and oleuropein, with respect to superoxide anion generation, neutrophils respiratory burst, and hypochlorous acid. low EC50s indicate that both compounds are potent scavengers of superoxide radicals and inhibitors of neutrophils respiratory burst: whenever demonstrated in vivo, these properties may partially explain the observed lower incidence of CHD and cancer associated with the Mediterranean diet.

FILE 'MARPAT' ENTERED AT 12:38:16 ON 03 MAR 2006
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FILE CONTENT: 1969-PRESENT (VOL 144 ISS 9 (20060224/ED)

SOME MARPAT RECORDS ARE DERIVED FROM INPI DATA FOR 1969-1987

MOST RECENT CITATIONS FOR PATENTS FROM FIVE MAJOR ISSUING AGENCIES (COVERAGE TO THESE DATES IS NOT COMPLETE):

US 2006014764 19 JAN 2006 DE 2020050148977 22 DEC 2005 EP 1605533 14 DEC 2005 JP 2005353222 22 DEC 2005 WO 2006003494 12 JAN 2006

Expanded G-group definition display now available.

New CAS Information Use Policies, enter HELP USAGETERMS for details.

L3 STR

VAR G1=0/S/CH2/19 VAR G2=0/S/CH2/19 VAR G3=OH/N/S NODE ATTRIBUTES: CONNECT IS X2 RC AT 15 CONNECT IS M3 RC AT 17 DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

ATTRIBUTES SPECIFIED AT SEARCH-TIME:

ECLEVEL IS LIM ON ALL NODES ALL RING(S) ARE ISOLATED

4 SEA FILE-MARPAT ABBON PLU-ON L17/COMPLETE - Retrieves only L18

7 SEA FILE=MARPAT SSS FUL L3 (MODIFIED ATTRIBUTES)

atations w

L18 ANSWER 1 OF 4 MARPAT COPYRIGHT 2006 ACS on STN iterations

ACCESSION NUMBER:

140:400046 MARPAT

TITLE:

L17

Methods for inhibiting cancer and scar formation

INVENTOR(S): Hamdi, Hamdi K.; Castellon, Raquel

PATENT ASSIGNEE(S):

SOURCE:

U.S. Pat. Appl. Publ., 24 pp., Cont.-in-part of

U.S. 657,414. CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004097428	A1	20040520	US 2003-712423	20031113
US 2003004117	A1	20030102	US 2002-153003	20020522
US 6632798	B2	20031014		
US 2004048808	A1	20040311	US 2003-657414	20030908
CA 2508786	AA	20040624	CA 2003-2508786	20031204
WO 2004053067	A2	20040624	WO 2003-US38564	20031204
WO 2004053067	A 3	20040819		
W: AE, AG,	AL, AM	, AT, AU, AZ,	BA, BB, BG, BR, BY	, BZ, CA, CH,
~~~	~~ ~	AR DE D21	D/ DA DA DA DA	70 PT 00

CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB,

571-272-2528 Searcher : Shears

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GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR,
             KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX,
             MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG,
             SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU,
             ZA, ZM, ZW
         RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
             AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE,
             DK, EE, ES, FI, FR, GB, GR, HU, IE, IT; LU, MC, NL, PT, RO,
             SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML,
             MR, NE, SN, TD, TG
                                            EP 2003-812800
                       A2 20050907
     EP 1569516
                                                               20031204
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
                                             US 2002-153003
PRIORITY APPLN. INFO.:
                                                               20020522
                                             US 2002-431780P 20021209
                                             US 2003-657414
                                                               20030908
                                             US 2001-292947P 20010523
                                                               20031113
                                             US 2003-712423
                                             WO 2003-US38564 20031204
AB
     Methods are disclosed for inhibiting cancer, scar formation,
     disrupting the cellular cytoskeleton, and conferring resistance from
     infection are disclosed. Such methods comprise the administration of
     oleuropein and/or the products of its hydrolysis in therapeutically
     effective amts. To that end, a variety of pharmaceutical formulations
     and routes or administration are disclosed and may be utilized to
```

L18 ANSWER 2 OF 4 MARPAT COPYRIGHT 2006 ACS on STN

treat a wide variety of diseases.

ACCESSION NUMBER:

140:229477 MARPAT

TITLE:

Methods using oleuropein and related compounds for

INVENTOR(S):

inhibiting angiogenesis, and therapeutic use Hamdi, Hamdi K.; Tavis, Jeffrey H.; Castellon,

Raquel

PATENT ASSIGNEE(S):

USA

SOURCE:

U.S. Pat. Appl. Publ., 23 pp., Cont.-in-part of

U.S. Ser. No. 153,003.

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO. DATE
US 2004048808	A1	20040311	US 2003-657414 20030908
US 2003004117	A1	20030102	US 2002-153003 20020522
US 6632798	B2	20031014	
US 2004097428	<b>A</b> 1	20040520	US 2003-712423 20031113
PRIORITY APPLN. INFO.	:		US 2001-292947P 20010523
			US 2002-153003 20020522
			US 2002-431780P 20021209
			US 2003-657414 20030908

Methods for inhibiting angiogensis are disclosed which comprise AB administering oleuropein and/or the products of its hydrolysis in therapeutically effective amts. The methods and compns. of the invention are particularly effective in inhibiting the vascularization of endothelial cells, and may be utilized to treat a wide variety of cancers, ocular diseases, and inflammatory conditions.

L18 ANSWER 3 OF 4 MARPAT COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 137:380057 MARPAT

TITLE: Methods for inhibiting angiogenesis using

oleuropein and its hydrolysis products

INVENTOR(S):
Hamdi, Hamdi K.; Tavis, Jeffrey H.; Castellon,

Raquel

PATENT ASSIGNEE(S): USA

SOURCE: PCT Int. Appl., 54 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

	PATENT NO.			KIND DATE				APPLICATION NO.					DATE				
	WO 2002094193			A1 20021128				WO 2002-US16191 2002052						0522			
		W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,
			CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,
			GΕ,	GH,	GM,	HR,	ΗU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	ΚZ,
			LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,
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AB Methods for inhibiting angiogenesis comprise administering oleuropein and/or the products of its hydrolysis in therapeutically effective amts. The methods and compns. of the present invention are particularly effective in inhibiting the vascularization of endothelial cells, and may be utilized to treat a wide variety of cancers, ocular diseases, and inflammatory conditions. For example, anti-angiogenic properties of oleuropein in the adult mouse ear model were illustrated. Oleuropein potently inhibited existing blood vessels from sprouting. The burn area is in fact devoid of blood vessels.

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L18 ANSWER 4 OF 4 MARPAT COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 125:96055 MARPAT

TITLE: Antiviral compositions comprising secoiridoids

obtained from Oleaceae

INVENTOR(S): Fredrickson, William R.
PATENT ASSIGNEE(S): Strecker, Robert B., USA
SOURCE: PCT Int. Appl., 20 pp.

CODEN: PIXXD2

DOCUMENT TYPE: LANGUAGE: Patent English

FAMILY ACC. NUM. COUNT:

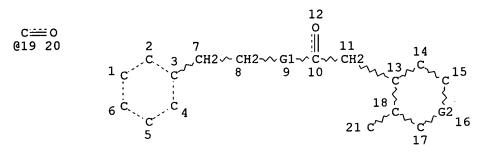
PATENT INFORMATION:

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WO	9614	064		Α	1	1996	0517		W	<b>19</b> :	95 <b>-</b> U	S140	56	1995	1103	
	W:	AL,	AM,	ΑŤ,	ΑU,	BB,	BG,	BR,	BY,	CA,	CH;	CN,	CZ,	DE,	DK,	EE,
		ES,	FI,	GB,	GE,	HU,	IS,	JP,	ΚE,	KG,	KP,	KR,	ΚZ,	LK,	LR,	LS,
		LT,	LU,	LV,	MD,	MG,	MK,	MN,	MW,	MX,	NO,	ΝZ,	PL,	PT,	RO,	RU,
		SD,	SE,	SG,	SI,	SK										
	RW:	ΚE,	LS,	MW,	SD,	SZ,	ŪG,	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,
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									U:	5 19	96-6	6832	4	1996	0626	

AB A method of treatment of diseases of viral origin comprises oral or parenteral administration of an antiviral amount of a naturally occurring secoiridoid or derivs. thereof from plants of the family Oleaceae. Preferred oral dosage forms include the secoiridoid oleuropein in pure form or as a component of dried plant material of Olea europaea or a dried extract thereof and a pharmaceutically acceptable carrier. Dried leaves of Olea europaea was suspended in 2 volume of red wine and held at room temperature for 7-10 days with periodic stirring. Filtration of the mixture provided a tincture containing 88 mg of oleuropein/oz of fluid. The antiviral efficacy of above composition in treatment of six subjects afflicted with herpes virus infection is reported.

FILE 'HOME' ENTERED AT 12:40:51 ON 03 MAR 2006

=> => d que stat 14; d que stat 118; d his ful L1 STR



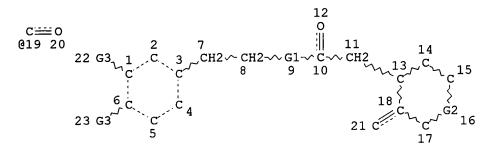
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GRAPH ATTRIBUTES:

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STEREO ATTRIBUTES: NONE

L2 ( 132) SEA FILE=REGISTRY SSS FUL L1
L3 STR



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VAR G3=OH/N/S
NODE ATTRIBUTES:
CONNECT IS X2 RC AT 15
CONNECT IS M3 RC AT 17
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

L4 32 SEA FILE=REGISTRY SUB=L2 SSS FUL L3

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VAR GI=0/5/CH2/19
VAR G2=0/5/CH2/19
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NODE ATTRIBUTES:
CONNECT IS X2 RC AT 15
CONNECT IS M3 RC AT 17
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES: RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

ATTRIBUTES SPECIFIED AT SEARCH-TIME: ECLEVEL IS LIM ON ALL NODES ALL RING(S) ARE ISOLATED

L17 7 SEA FILE=MARPAT SSS FUL L3 (MODIFIED ATTRIBUTES)
L18 4 SEA FILE=MARPAT ABB=ON PLU=ON L17/COMPLETE

(FILE 'REGISTRY' ENTERED AT 12:27:31 ON 03 MAR 2006)
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L1 STR
L2 ( 132)SEA SSS FUL L1
L3 STR
L4 32 SEA SUB=L2 SSS FUL L3

FILE 'REGISTRY' ENTERED AT 12:30:53 ON 03 MAR 2006 D QUE STAT

FILE 'CAPLUS' ENTERED AT 12:30:53 ON 03 MAR 2006
L5 540 SEA ABB=ON PLU=ON L4
L6 30 SEA ABB=ON PLU=ON L5 AND (?CANCER? OR ?CARCIN? OR ?TUMOUR? OR ?TUMOR? OR ?NEOPLAS?)
L7 13 SEA ABB=ON PLU=ON L6 NOT (PY=>2002 OR PD=>20021209)

L*** DEL 4 S L6 AND HAMDI ?/AU

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FILE 'REGISTRY' ENTERED AT 12:31:55 ON 03 MAR 2006 L*** DEL 4 S E30-33

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SEL HIT L8 2 RN

FILE 'CAPLUS' ENTERED AT 12:33:43 ON 03 MAR 2006 SEL HIT L7 1-13 RN D L7 1-13 IBIB ABS HITSTR

FILE 'USPATFULL' ENTERED AT 12:34:48 ON 03 MAR 2006
L9 74 SEA ABB=ON PLU=ON L4

L10 27 SEA ABB=ON PLU=ON L9 AND (?CANCER? OR ?CARCIN? OR ?TUMOUR? OR ?TUMOR? OR ?NEOPLAS?)

L11 3 SEA ABB=ON PLU=ON L10 NOT (PY=>2002 OR PD=>20021209) D 1-3 IBIB ABS

FILE 'MEDLINE, BIOSIS, EMBASE' ENTERED AT 12:35:57 ON 03 MAR 2006 L*** DEL 406 S L4

FILE 'MEDLINE, BIOSIS, EMBASE' ENTERED AT 12:36:35 ON 03 MAR 2006 L12 406 SEA ABB=ON PLU=ON L4

L12 406 SEA ABB=ON PLU=ON L4
L13 50 SEA ABB=ON PLU=ON L12 AND (CANCER? OR CARCIN? OR TUMOUR?
OR TUMOR? OR NEOPLAS? OR ANTICANCER? OR ANTICARCIN? OR
ANTITUMOUR? OR ANTITUMOR? OR ANTINEOPLAS?)

L14 35 DUP REM L13 (15 DUPLICATES REMOVED)
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L15 0 SEA ABB=ON PLU=ON L12(L)(CANCER? OR CARCIN? OR TUMOUR?
OR TUMOR? OR NEOPLAS? OR ANTICANCER? OR ANTICARCIN? OR
ANTITUMOUR? OR ANTITUMOR? OR ANTINEOPLAS?)
D L14 1-35 IBIB ABS

FILE 'MARPAT' ENTERED AT 12:38:16 ON 03 MAR 2006 D QUE L3

L16 0 SEA SSS SAM L3 (MODIFIED ATTRIBUTES)

L17 7 SEA SSS FUL L3 (MODIFIED ATTRIBUTES)

L18 4 SEA ABB=ON PLU=ON L17/COMPLETE

D QUE STAT D 1-4 .BEVMAR1

FILE 'HOME' ENTERED AT 12:40:51 ON 03 MAR 2006
D QUE STAT L4
D QUE STAT L18

FILE REGISTRY

Property values tagged with IC are from the ZIC/VINITI data file

provided by InfoChem.

STRUCTURE FILE UPDATES: 2 MAR 2006 HIGHEST RN 875740-40-2 DICTIONARY FILE UPDATES: 2 MAR 2006 HIGHEST RN 875740-40-2

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* The CA roles and document type information have been removed from * the IDE default display format and the ED field has been added, * effective March 20, 2005. A new display format, IDERL, is now * available and contains the CA role and document type information. * *

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http://www.cas.org/ONLINE/UG/regprops.html

FILE CAPLUS

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FILE CAOLD
FILE COVERS 1907-1966
FILE LAST UPDATED: 01 May 1997 (19970501/UP)

This file contains CAS Registry Numbers for easy and accurate substance identification. Title keywords, authors, patent assignees, and patent information, e.g., patent numbers, are now searchable from 1907-1966. TIFF images of CA abstracts

printed between 1907-1966 are available in the PAGE display formats.

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FILE USPATFULL

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FILE COVERS 1971 TO PATENT PUBLICATION DATE: 2 Mar 2006 (20060302/PD)
FILE LAST UPDATED: 2 Mar 2006 (20060302/ED)
HIGHEST GRANTED PATENT NUMBER: US7007305
HIGHEST APPLICATION PUBLICATION NUMBER: US2006048257
CA INDEXING IS CURRENT THROUGH 28 Feb 2006 (20060228/UPCA)
ISSUE CLASS FIELDS (/INCL) CURRENT THROUGH: 2 Mar 2006 (20060302/PD)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Dec 2005
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Dec 2005

# FILE MEDLINE

FILE LAST UPDATED: 2 MAR 2006 (20060302/UP). FILE COVERS 1950 TO DAT

On December 11, 2005, the 2006 MeSH terms were loaded.

The MEDLINE reload for 2006 is now (26 Feb.) available. For details on the 2006 reload, enter HELP RLOAD at an arrow prompt (=>). See also:

http://www.nlm.nih.gov/mesh/

http://www.nlm.nih.gov/pubs/techbull/nd04/nd04 mesh.html

http://www.nlm.nih.gov/pubs/techbull/nd05/nd05 med data changes.ht

http://www.nlm.nih.gov/pubs/techbull/nd05/nd05 2006 MeSH.html

OLDMEDLINE is covered back to 1950.

MEDLINE thesauri in the /CN, /CT, and /MN fields incorporate the MeSH 2006 vocabulary.

This file contains CAS Registry Numbers for easy and accurate substance identification.

FILE BIOSIS

FILE COVERS 1969 TO DATE.

CAS REGISTRY NUMBERS AND CHEMICAL NAMES (CNs) PRESENT FROM JANUARY 1969 TO DATE.

RECORDS LAST ADDED: 1 March 2006 (20060301/ED)

# FILE EMBASE

FILE COVERS 1974 TO 24 Feb 2006 (20060224/ED)

EMBASE has been reloaded. Enter HELP RLOAD for details.

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FILE MARPAT

FILE CONTENT: 1969-PRESENT (VOL 144 ISS 9 (20060224/ED)

SOME MARPAT RECORDS ARE DERIVED FROM INPI DATA FOR 1969-1987

MOST RECENT CITATIONS FOR PATENTS FROM FIVE MAJOR ISSUING AGENCIES (COVERAGE TO THESE DATES IS NOT COMPLETE):

US 2006014764 19 JAN 2006
DE 2020050148977 22 DEC 2005
EP 1605533 14 DEC 2005
JP 2005353222 22 DEC 2005
WO 2006003494 12 JAN 2006

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